



Geochemical Data for Core and Bottom-Sediment Samples Collected in 2007 from Grand Lake O' the Cherokees, Northeast Oklahoma

By David L. Fey, Mark F. Becker, and Kathleen S. Smith

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Conversion Factors

Inch/Pound to SI

Multiply	By	To obtain
Length		
inch (in)	2.54	centimeter (cm)
inch (in)	0.0254	meter (m)
foot (ft)	0.3048	meter (m)
mile (mi)	1.609	kilometer (km)
Area		
square mile (mi^2)	2.590	square kilometer (km^2)
Mass		
ounce, avoirdupois (oz)	28.35	gram (g)
pound, avoirdupois (lb)	453.6	gram (g)
ton, short (2,000 lb)	0.9072	megagram (Mg)

SI to Inch/Pound

Multiply	By	To obtain
Length		
centimeter (cm)	0.3937	inch (in)
meter (m)	39.37	inch (in)
meter (m)	3.281	foot (ft)
kilometer (km)	0.6214	mile (mi)
Area		
square kilometer (km^2)	0.3861	square mile (mi^2)
Mass		
gram (g)	0.03527	ounce, avoirdupois (oz)
gram (g)	0.002205	pound avoirdupois (lb)
megagram (Mg)	1.102	ton, short (2,000 lb)

Temperature in degrees Celsius ($^{\circ}\text{C}$) may be converted to degrees Fahrenheit ($^{\circ}\text{F}$) as follows:

$$^{\circ}\text{F} = (1.8 \times ^{\circ}\text{C}) + 32$$

Temperature in degrees Fahrenheit ($^{\circ}\text{F}$) may be converted to degrees Celsius ($^{\circ}\text{C}$) as follows:

$$^{\circ}\text{C} = (^{\circ}\text{F} - 32) / 1.8$$

Vertical coordinate information is referenced to the World Geodetic System of 1984 (WGS84).

Horizontal coordinate information is referenced to the World Geodetic System of 1984 (WGS84).

Altitude, as used in this report, refers to distance above the vertical datum.

Concentrations of chemical constituents in solid samples are given in either percent (%), parts per million (ppm), or parts per billion (ppb).

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Introduction

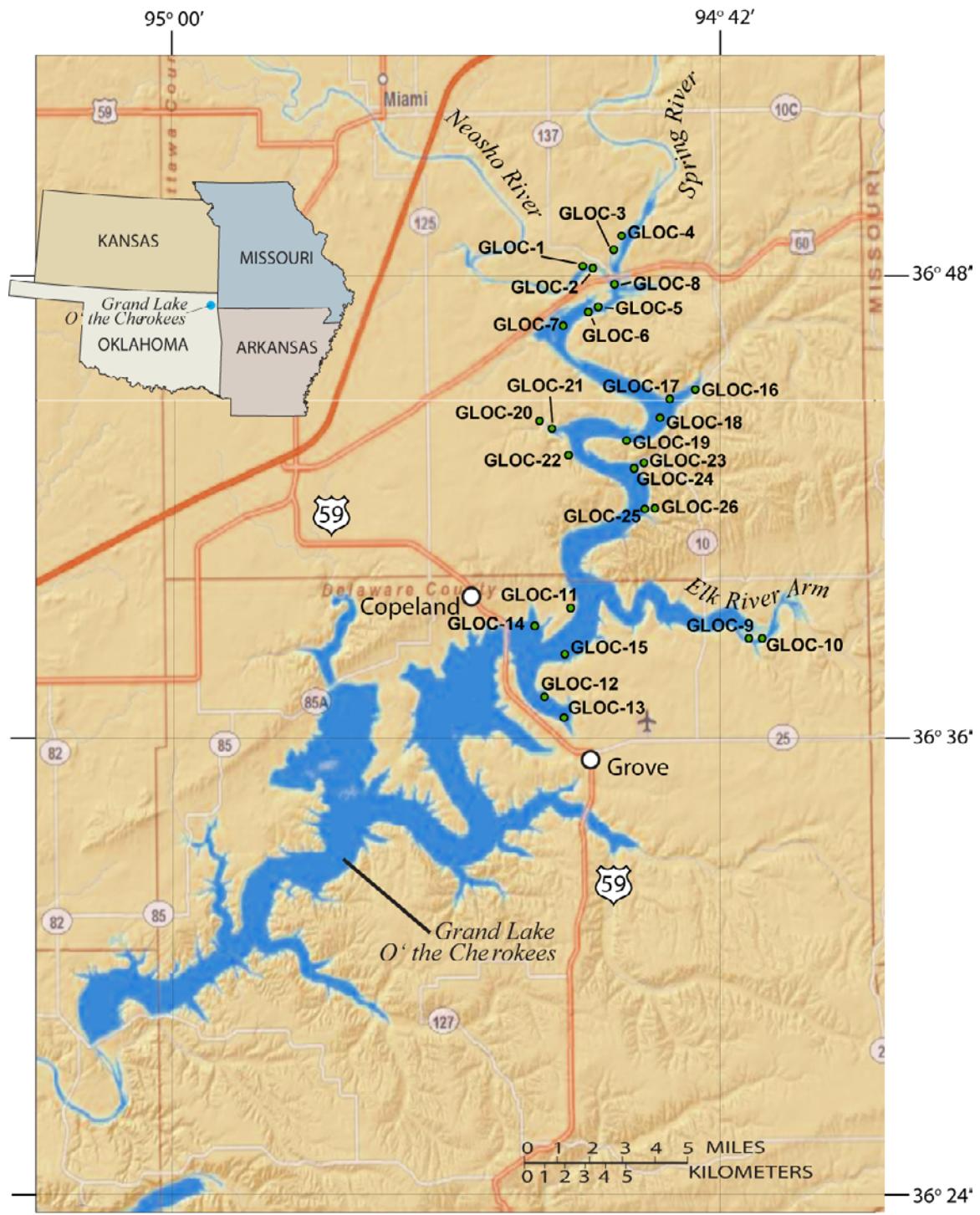
The Tri-State Mining District, a Mississippi Valley-type (MVT) zinc-lead district, covers an area of about 1,200 square miles (mi^2) (3,108 square kilometers [km^2]) in northeast Oklahoma, southwest Missouri, and southeast Kansas. The district was active from about 1850 to 1970 and during much of this time was the largest producer of zinc (Zn) and lead (Pb) ore in the world. Over the lifespan of the district, more than 4,000 mines produced 23 million tons of zinc concentrate and 4 million tons of lead concentrate (Brosius and Sawin, 2001). Gibson (1982) estimated the district total production at 8.8 million tons zinc and 1.7 million tons lead.

A consequence of 120 years of mining activity in the district was the distribution on the surface of between 165 and 300 million tons of processed mine tailings (chat), the majority of which has been used as road aggregate or has eroded into nearby drainages. The U.S. Army Corps of Engineers has estimated that 75 million tons of chat remain in the district (Andrews and others, 2009; State of Oklahoma, 2000). This material contains elevated concentrations of cadmium (Cd), lead, and zinc (Andrews and others, 2009; Juracek and Becker, 2009), which can be transported by water and wind into nearby receiving streams, ponds, and rivers. Seepage from flooded mine workings and the leaching of chat material can also transport dissolved metals to local surface water (Andrews and others, 2009). The metals are transported either as dissolved aqueous species downstream (Carroll and others, 1998), as mineral grains entrained in bottom sediment, or through sorption to colloidal material or clay that becomes incorporated into the bottom sediment of streams, rivers, lakes, and reservoirs (Andrews and others, 2009). Accumulation of these metals in sediments downstream from the district is recognized as a potential hazard to aquatic life (Pope, 2005; Andrews and others, 2009).

Grand Lake O' the Cherokees is a large reservoir in northeast Oklahoma, below the confluence of the Neosho and Spring Rivers, both of which drain the Tri-State District to the north (fig. 1). The reservoir was constructed in 1940 by the completion of the Pensacola Dam by the Grand River Dam Authority. Concerns of sediment quality and the possibility of human exposure to cadmium and lead through eating fish have led to several studies of the sediments in the Tri-State District:

- bottom sediment of Grand Lake O' the Cherokees: Juracek and Becker (2009);
- stream sediment in the Oklahoma part of the Tri-State District: Andrews and others (2009);
- Empire Lake, Kansas: Juracek (2006);
- stream sediment in the Kansas part of Tri-State district: Pope (2005);
- stream sediment in the Tar Creek Basin: Parkhurst and others (1988); and
- stream sediment in the Tri-State District: Carroll and others (1998).

Other references regarding the Tar Creek and Picher areas include Erickson and others (1988), Parkhurst (1987, 1988), Parkhurst and others (1988), and Smith and others (1988).



Base map from ArcGIS online services, ESRI, 2009, Redlands Calif.
Inset map from Juracek and Becker (2009).

Figure 1. Location map for samples taken in the Grand Lake O' the Cherokees area in northeast Oklahoma.

In order to record the transport and deposition from the Tri-State District of metals by the Spring and Neosho Rivers, the U.S. Geological Survey (USGS) collected 11 sediment cores and 15 bottom-sediment samples from the Grand Lake O' the Cherokees in northeastern Oklahoma in September 2007. Subsamples from five selected cores and the bottom-sediment samples were analyzed for major and trace elements and forms of carbon. Presentation of the analytical results is the purpose of this report.

Methods

Sediment Core Collection and Preparation

Cores of reservoir sediment were collected at 11 sites from the Grand Lake O' the Cherokees reservoir (fig. 1). These cores were labeled GLOC-1 (Grand Lake O' the Cherokees) through GLOC-11. A 3-inch-diameter aluminum pipe (barrel) was driven down into the sediment from a pontoon raft working platform (fig. 2). A gasoline engine-powered vibrator (typically used for settling wet concrete) was attached to the barrel; this arrangement allowed the pipe to be driven to depths of 0.8–3 meters (m) (31–120 inches [in]) below the sediment-water interface (fig. 3). The depth of water where cores were taken varied from 0.6 to 3 m (2 to 10 feet [ft]). Attempts to work in deeper water were made but were unsuccessful.



Figure 2. Pontoon boat coring platform. Aluminum tower is for attaching hand-operated winch to hoist core barrel.



Figure 3. Close-up view of mechanical vibrator attached to core barrel. Core is standing through hole in platform deck; gasoline engine in background is used to drive the vibrator.

After the core barrel was driven to refusal, the barrel was extracted from the sediment using a hand-powered hoist attached to the tower on the working platform. When possible, the depth of penetration was recorded. The bottom of the core barrel was capped as it was lifted above the water surface. Each core barrel was transported vertically to the shore, where the position of the top of the sediment was determined, yielding the core recovery length. The remaining barrel above was cut away, and the top of the recovered core was packed with laboratory cleaning tissue (to prevent mixing of the top of core) and capped. Cores were transported to a USGS workstation warehouse in Oklahoma City, Okla. This warehouse is located at a locked, secure General Services Administration (GSA) compound. Keys to the warehouse are accessible to USGS Oklahoma Water Science Center personnel only.

The cores were stored in an upright position in a freezer from September 2007 to July 2008, at which time they were processed to yield subsamples. The cores remained sealed and frozen for the entire period of storage; taped end caps were undisturbed. In the warehouse, each core was laid horizontally in a jig, and the aluminum barrel was cut longitudinally its entire length with a hand-held circular saw. The blade depth of the saw was set to just penetrate the aluminum wall in order to avoid incorporating possible metal fragments into the sediment. The core was then rotated 180 degrees and the other side was cut in the same manner. One cut was taped with duct tape to form a hinge. The cored sample material was then cut with a plastic knife longitudinally, and one side of the barrel was rotated to expose both halves of the core. Each core was subsampled at 5-centimeter (cm) (2-in) intervals. Care was taken not to allow any contamination from the aluminum core barrel: the exterior of each subsample was scraped off and the interior sediment was carefully removed and placed in a clean glass bowl to air dry in a closed hood. A subsample of the core was taken from each interval for chemical analysis, and another subsample was taken from core GLOC-7 for ^{137}Cs (cesium-137) dating. Mark Becker of the USGS Oklahoma Water Science Center performed these tasks.

The subsamples for chemical analysis and ^{137}Cs dating were sent by express mail to Kathleen Smith and David Fey at the USGS laboratories in Building 20 at the Denver Federal Center (DFC) in Lakewood, Colo. Building 20 is a secure building, accessible by proximity card by authorized USGS personnel only. The Denver Federal Center is a limited-access facility with guarded entrances.

At the DFC labs, the samples were visually described, sieved to minus-80 mesh (177 micrometers [μm]), and ground to minus-100 mesh (150 μm). The ground samples were then split, with one set of splits kept for archival storage at the DFC and the other arranged into lots for submittal to the USGS contract laboratory SGS Minerals Services in Toronto, Canada, for analysis. Mineralogical analysis by X-ray diffraction (XRD) was performed at the Denver USGS laboratory on selected sieved, unground subsamples from cores GLOC-2, GLOC-3, and GLOC-7. During the sample description, preparation, XRD determinations, and shipping processes, the samples were kept in locked offices, locked labs, or the secure Sample Control facility contained within Building 20.

Bottom Sediment Collection and Preparation

Collecting core samples was time consuming and, in addition, not possible in deeper water, so we supplemented the sediment cores with reservoir bottom-sediment samples collected with a simple gravity corer. A single composite sample of reservoir bottom sediment was obtained at 15 sites by dropping a simple 9-cm-diameter (3.5-in-diameter) iron gravity corer from the working platform (fig. 4). The corer was unlined, and no attempt was made to preserve the stratigraphy of the samples; a composite sample was made from the sediment collected from about the top 18 cm (7 in) of bottom sediment. These cores were labeled GLOC-12 through GLOC-26. After collection, the bottom-sediment samples were double-bagged in large plastic storage bags and transported to the USGS Oklahoma Water Science Center warehouse described above. The bottom-sediment samples remained frozen during storage, from September 2007 to July 2008.



Figure 4. Simple gravity corer used to collect reservoir bottom-sediment samples.

Samples were air-dried at the warehouse and then shipped to the Denver USGS laboratories for preparation. They were processed in the same manner as the core subsamples: sieved to minus-80 mesh, ground to minus-100 mesh, arranged in lots, and sent to the USGS contract laboratory SGS Minerals Services in Toronto, Canada.

Sample Localities and Descriptions

Sample localities are plotted in figure 1 and listed in table 1. Table 2 contains depths of water and descriptions for the core and bottom-sediment samples.

Table 1. Field numbers, sample localities, dates, and site descriptions for core and reservoir bottom-sediment samples taken from Grand Lake O' the Cherokees, September 2007. ([Click to see Excel file.](#))

[Sample coordinates are referenced to the World Geodetic System of 1984 and are given in decimal degree. USF, upstream from; DSF, downstream from; mi, mile; Hwy, highway]

Core number	North latitude	West longitude	Date	Site description
GLOC-1	36.80410	94.76656	9/12/2007	Neosho River about 0.7 mi USF Hwy 60 bridge
GLOC-2	36.80331	94.76100	9/12/2007	Neosho River about 0.4 mi USF Hwy 60 bridge
GLOC-3	36.81125	94.74951	9/13/2007	Spring River about 0.8 mi USF Hwy 60 bridge
GLOC-4	36.81717	94.74482	9/13/2007	Spring River about 1.3 mi USF Hwy 60 bridge
GLOC-5	36.78646	94.75837	9/13/2007	reservoir about 0.5 mi DSF railroad bridge
GLOC-6	36.78439	94.76396	9/13/2007	reservoir about 0.3 mi DSF from GLOC-5
GLOC-7	36.777851	94.77760	9/13/2007	reservoir about 0.9 mi DSF from GLOC-6
GLOC-8	36.79668	94.75351	9/13/2007	reservoir at confluence of Spring and Neosho Rivers
GLOC-9	36.64159	94.68297	9/14/2007	Elk River about 1.7 mi USF from Cowskin Bridge
GLOC-10	36.64157	94.67565	9/14/2007	Elk River about 2.1 mi USF from Cowskin Bridge
GLOC-11	36.65591	94.77567	9/14/2007	reservoir right bank, across from Elk River arm
GLOC-12	36.61737	94.79456	9/15/2007	Wolf Creek Cove
GLOC-13	36.60817	94.78305	9/15/2007	upper Wolf Creek Cove
GLOC-14	36.64812	94.79848	9/15/2007	reservoir near Hickory Creek Island
GLOC-15	36.63576	94.77902	9/15/2007	reservoir main channel near Walnut Cove
GLOC-16	36.74973	94.71201	9/15/2007	upper Sycamore Cove
GLOC-17	36.74627	94.72341	9/15/2007	lower Sycamore Cove
GLOC-18	36.74020	94.72669	9/15/2007	mouth of Sycamore Cove
GLOC-19	36.72820	94.74387	9/15/2007	reservoir between Sycamore and Bee Creek Coves
GLOC-20	36.73712	94.79139	9/15/2007	upper Bee Creek Cove
GLOC-21	36.73369	94.78447	9/15/2007	mouth of Bee Creek Cove
GLOC-22	36.72219	94.77583	9/15/2007	reservoir about 0.9 mi below Bee Creek Cove
GLOC-23	36.71917	94.73907	9/15/2007	upper Council Cove
GLOC-24	36.71709	94.74388	9/15/2007	mouth of Council Cove
GLOC-25	36.69832	94.73851	9/15/2007	main channel, near North Broad Hollow
GLOC-26	36.69864	94.73283	9/15/2007	upper North Broad Hollow

Table 2. Field numbers, water depths, and visual sample descriptions for core and reservoir bottom-sediment samples taken from Grand Lake O' the Cherokees, September 2007. ([Click to see Excel file.](#))

[m, meter; in, inch]

Core number	Depth of water (m)	Sample description
GLOC-1	3	core: silt and clay, sparse fine sand
GLOC-2	0.6	core: silt and clay, sparse fine sand
GLOC-3	0.9	core: fine sand and silt, little clay
GLOC-4	0.6	core: brown/gray clay, some silt at bottom; fine brown sand
GLOC-5	1.2	core: 63 in penetration; 38 in recovery
GLOC-6	6.7	no sample taken; encountered bedrock at bottom of reservoir
GLOC-7	1.2	core: gray clayey silt; 72 in penetration, 54 in recovery
GLOC-8	1.2	core: penetration/recovery not recorded
GLOC-9	1.2	core: 94 in penetration, 90 in recovery
GLOC-10	1.2	core: 120 in penetration, 90 in recovery; brown clay
GLOC-11	1.5	core: 32 in penetration, recovery not recorded
GLOC-12	5.2	reservoir sediment: dark gray very fine sand
GLOC-13	2.4	reservoir sediment: dark gray very fine sand
GLOC-14	6.7	reservoir sediment: dark gray very fine sand
GLOC-15	13	reservoir sediment: dark gray very fine sand
GLOC-16	2.1	reservoir sediment: dark gray very fine sand
GLOC-17	2.1	reservoir sediment: dark gray very fine sand with quartz grains
GLOC-18	1.5	reservoir sediment: gray very fine sand
GLOC-19	6.7	reservoir sediment: gray very fine sand
GLOC-20	0.9	reservoir sediment: gray very fine sand
GLOC-21	4.3	reservoir sediment: gray very fine sand
GLOC-22	7.3	reservoir sediment: gray very fine sand
GLOC-23	2.4	reservoir sediment: gray-brown very fine sand
GLOC-24	4.3	reservoir sediment: gray-brown very fine sand
GLOC-25	7.9	reservoir sediment: gray-brown very fine sand
GLOC-26	4.9	reservoir sediment: gray-brown very fine sand

X-Ray Diffraction Analyses

Forty-one selected core subsamples were analyzed for mineralogical content by X-ray diffraction (XRD) using a Shimadzu XRD-6000 X-Ray diffractometer equipped with a standard copper (Cu) K-alpha radiation tube, monochromometer, and automatic 5-sample changer. All samples were powdered using an agate mortar and pestle. Each sample powder was mixed with a few drops of acetone to form a slurry, which was transferred by pipette to a zero-background quartz slide. These slurry mounts were inserted into the sequential auto sample changer.

Raw XRD data acquired from the instrument were analyzed using Jade Search/Match software, version 7.5. The data were converted to diffractograms, which were background-reduced and smoothed. Mineral phase identities, selected from the International Centre for Diffraction Data (ICDD) database, were assigned to all visible peaks. Phase quantification was based on a visual comparison of peak intensities and assigned *M* for major constituent, *m* for minor constituent, and *t* for trace values. Inconclusive phase identifications were assigned a *p* for possible. Table 3 presents the XRD data for a subset of samples from cores GLOC-2, GLOC-3, and GLOC-7.

Table 3. Field numbers and mineralogy by X-ray diffraction for subsamples from cores GLOC-2, GLOC-3, and GLOC-7 from Grand Lake O' the Cherokees (GLOC). No sulfide minerals were detected by this method.
[\(Click to see Excel file.\)](#)

[cm, centimeter; M, major constituent; m, minor constituent; p, possible, followed by either m or t; t,trace. Qtz, quartz; Ms,muscovite; Ill,Illite; Phg,phengite; Ab,albite; Mc,microcline; Or,orthoclase; Sa,sanidine; Kln,Kaolinite; Chl,chlorite; Sme, smectite]

Core number	Depth below water/sediment interface (cm)	Qtz	Ms/Ill	Ms/Phg	Ab	Mc	Or/Sa	Kln/Chl	Sme
GLOC-2 1-3	5	M	m		m		m		
GLOC-2 5-7	15	M	m		m	m		p,t	p,t
GLOC-2 9-11	25	M	m		m	m		p,t	
GLOC-2 13-15	35	M	m		m	m		p,t	p,t
GLOC-2 17-19	45	M	m		m			p,t	p,t
GLOC-2 21-23	55	M	m		m	m	m		t
GLOC-2 25-27	65	M	m		m		m		p,t
GLOC-2 29-31	75	M	m		m	m		p,t	
GLOC-2 33-35	85	M	m		m		m		m
GLOC-2 37-39	95	M	m		m	m			
GLOC-2 41-43	105	M	m		m	m			
GLOC-2 45-47	115	M	m		m	m	m	p,t	m
GLOC-2 55-57	140	M	m		m	m			p,t
GLOC-2 65-67	165	M	m		m	m			p,t
GLOC-2 75-77	190	M	m		m		m	t	t
GLOC-2 85-87	215	M	m			m	m		
GLOC-2 95-97	240	M	M			m	m		
GLOC-2 105-107	265	M	M			m	m		
GLOC-2 115-117	290	M	m		m				
GLOC-2 121-123	300	M	m		m		m	p,t	p,t
GLOC-3 0-2	2.5	M	m		p,t				
GLOC-3 4-6	13	M		m	m	m			
GLOC-3 8-10	23	M	t		t	t			
GLOC-3 12-14	33	M	t		m	m	m	p,t	
GLOC-3 16-18	38	M	t		m		m		
GLOC-3 20-22	53	M	m		m	m			
GLOC-3 24-26	63	M		t	m	m			
GLOC-3 28-30	73	M	m			m	m		p,t
GLOC-3 32-34	83	M	t		m	p,m		p,t	
GLOC-3 36-38	93	M	m		m	m			m
GLOC-3 40-42	103	M	t				m		p,t
GLOC-3 44-46	113	M	m		m	m			
GLOC-3 48-50	123	M	m		m		m		
GLOC-3 52-54	133	M	m		m	m			p,t
GLOC-3 56-58	143	M	t		m	m	m		p,t
GLOC-3 60-62	153	M		t	t	t			p,t
GLOC-3 64-66	163	M	t				m		
GLOC-3 68-70	173	M	t		t	t			
GLOC-3 72-74	178	M	t		m	m	m		
GLOC-7 6-8	18	M	m		M		M	p,t	p,t
GLOC-7 38-40	98	M		m	m		m		

Multi-Element Analysis

Processed core subsamples and reservoir bottom sediment samples were weighed at SGS Mineral Services laboratories into 0.2-gram (g) aliquots and digested in a mixed four-acid medium of hydrochloric, nitric, perchloric, and hydrofluoric acids (HCl, HNO₃, HClO₄, and HF, respectively). The resulting solutions were analyzed for 26 elements by both inductively coupled plasma–atomic emission spectroscopy (ICP-AES) (Briggs, 2002) and inductively coupled plasma–mass spectrometry (ICP-MS) (Briggs and Meier, 2002). The major elements are reported from the ICP-AES instrument, whereas the trace elements are reported from either the ICP-AES or ICP-MS instrument. Instrument sensitivities, linear range, and freedom from spectral or mass interferences determine which instrument is used to report which elements. Table 1–1 in the appendix lists the elements analyzed, their lower reporting limits, and, for the 26-element package, which instrument was used for each element (ICP-AES or ICP-MS). The digestion procedure dissolves most minerals, including silicates, oxides, and sulfides; resistant or refractory minerals such as zircon, chromite, some rare-earth minerals, and some tin oxides are only partially dissolved. Previous investigations using a variety of geologic materials confirm the completeness of the digestion (Church and others, 1987; Wilson and others, 1994). For the purposes of analyzing for Mississippi Valley–type mineral deposit-related elements, the digestion is well suited.

Forms of Carbon Analyses

The core subsamples and bottom-sediment samples were analyzed for total carbon and carbonate carbon. The organic carbon content is obtained by the difference between those constituents. The total carbon was determined by combustion (Brown and Curry, 2002), where a 0.2-g sample is heated to 1,350°C to convert all forms of carbon to carbon dioxide, which is then measured using an infrared detector. The carbonate carbon was determined by a coulometric titration (Brown and others, 2002), where the sample is treated with hot 2 Normal perchloric acid and the evolved carbon dioxide is passed through a solution of monoethanolamine, into which it is quantitatively absorbed. This solution is then coulometrically titrated. The lower reporting limit for organic carbon is 0.05 percent.

Results

The analytical results for five of the sediment cores and the 15 bottom-sediment samples are in tables 1–2 through 1–7 in appendix 1. In this section are brief discussions with descriptive statistics for each table. To aid in these discussions, table 4 provides estimated crustal abundance values (Fortescue, 1992) for all constituents analyzed.

Table 4. Crustal abundance values for elements analyzed in this report. ([Click to see Excel file.](#))

[Values taken from Fortescue (1992). ppm, parts per million]

Element	Crustal abundance	Unit
Al	8.36	percent
Ca	4.66	percent
Fe	6.22	percent
K	1.84	percent
Mg	2.76	percent
Na	2.27	percent
P	0.112	percent
Ti	0.632	percent
S	340	ppm
As	1.8	ppm
Ba	390	ppm
Bi*	0.0082	ppm
Cd	0.16	ppm
Co	29	ppm
Cr	122	ppm
Cu	68	ppm
Ga	19	ppm
In	0.24	ppm
Li	18	ppm
Mn	1,060	ppm
Ni	99	ppm
Pb	13	ppm
Rb	78	ppm
Sr	384	ppm
V	136	ppm
Zn	76	ppm

*Value for Bi commented by Fortescue to be too small.

Bottom-Sediment Samples (Table 1–2)

The 15 bottom-sediment samples were taken from throughout the Grand Lake O' the Cherokees reservoir above Highway 59 bridge between Grove and Copeland and from several arms of the reservoir (samples GLOC-12 through GLOC-26 on fig. 1). These samples were collected by gravity corer and represent about the top 18 cm (7 in) of sediment. The top three intervals together of each sediment core also represent about 18 cm. In table 1–2, a ‘composite’ analysis has been made of the analyses of the first three intervals from cores GLOC-2, GLOC-3, GLOC-7, GLOC-8, and GLOC-9. The inclusion of these composite analyses increases the number of points in the reservoir with metals-content analyses of the bottom sediments at the sediment-water interface.

The XRD determinations for sediment mineralogy (table 3) show that the major constituent for all samples from the bottom surface and the cores is quartz. Predominance of quartz in the samples is also supported by the major-element analyses in table 1–2. By converting the major-element analyses to their respective oxide forms, summing them, and subtracting that sum from 100 percent, a rough estimate can be made of the SiO₂ content of the sediments. For the bottom-sediment samples in table 1–2, the average of this estimated SiO₂ content is 80 percent quartz equivalent. Some of the silica is contained in clays.

Other elements of table 1–2 are depleted compared to crustal abundance, again reflecting that the stream/river/lake environment has favored the retention of SiO₂. Evaluating the calcium analyses shows that, on average, the ratio of calcium to crustal abundance is only 18 percent. This deficiency of calcium reflects the fact that although much of the upstream rock is carbonate, it dissolves and releases calcium into solution. Other major elements with low ratios to crustal abundance are magnesium (Mg) and sodium (Na; each 17 percent), iron (Fe; 43 percent), and titanium (Ti; 38 percent). The magnesium has probably been mobilized like the calcium, whereas the low numbers for Na, Fe, and Ti probably reflect low concentrations in the original carbonate rocks.

Three major elements show higher ratios to crustal abundance. Aluminum (Al) has a value of 70 percent, and potassium (K) has a value of 77 percent, reflecting the clay content in the sediment. Sulfur (S) had a ratio to crustal abundance of 310 percent; although the highest concentration of sulfur was only 0.33 percent, the high ratio may be due to runoff from the metallic sulfide mining districts upstream of the reservoir.

The trace elements had ratios to crustal abundance mostly near or below 1.0, with exceptions being Zn (7.2), Cd (21), As (arsenic; 4.5), and Pb (3.3). The crustal abundance for zinc is 76 parts per million (ppm); the concentrations in the bottom-sediment samples ranged from 56 ppm (background, GLOC-9, Elk River Arm, table 1–7) to 1,523 ppm (20 times crustal abundance; GLOC-8, confluence of Neosho and Spring Rivers, table 1–6). Zinc is the most likely indicator of input from the mining districts, detected in sediment samples collected upstream of the reservoir in both the Spring and Neosho Rivers, and plays a major role in the discussions of the sediment cores. The zinc concentration at the top of core GLOC-2 (the surface sediment) collected in the Neosho River from the reservoir was only 122 ppm, less than 2 times crustal abundance. This enrichment is relatively slight, given that this location is downstream from the Picher district and Tar Creek (Andrews and others, 2009). The zinc concentration at the top of core GLOC-3 was 1,370 ppm. This relative enrichment of zinc indicates that sediment from the Spring River predominated the deposition at the surface at site GLOC-8, just below the confluence of the Spring and Neosho Rivers. Andrews and others (2009) noted that dissolved zinc concentrations in the Spring River might be higher than those in the Neosho River because a larger proportion of the Spring River basin had been mined (1.9 percent) than of the Neosho River basin (0.34 percent).

Overall, the distribution of zinc in the reservoir bottom sediments shows a general elevation of zinc in those drainages affected by historical mining operations. Of the 20 bottom-sediment samples represented in table 1–2 (15 gravity corer samples and 5 core upper interval composite analyses), 13 averaged 490 ppm, with only a 15 percent standard deviation. Thus, the zinc is fairly evenly distributed at the surface of the sediment. Outlier numbers can be explained by contribution from local terrestrial sediment input. For example, site GLOC-20 is located at the head of Bee Cove, and the sample was collected within meters of where Bee Creek enters the reservoir. The zinc concentration of that sample was 242 ppm. The local terrestrial sedimentation would serve to dilute, for a short distance, the effect of higher zinc concentrations in the sediment of the reservoir. Sample GLOC-21 supports this hypothesis. Located near the mouth of Bee Cove, just 0.8 km (0.5 mi) away, but exposed to the reservoir, the zinc

concentration of this sample was 528 ppm. Likewise, GLOC-23 is located at the upper end of Council Cove and contained only 134 ppm zinc. Local sedimentation also may be diluting the cove sediment at that location. Just 0.5 km away (0.3 mi) at the mouth of Council Cove, sample GLOC-24 contained 478 ppm zinc. As mentioned above, a sample was taken from the upper part of the Elk River arm of the reservoir (core GLOC-9). The Elk River drains a watershed with no known mining activity, and so sediment collected from that arm serves as a background comparison. The average zinc concentration from the top three intervals of GLOC-9 was only 56 ppm.

Cadmium, which is chemically similar to zinc and substitutes in the sphalerite lattice, had a high average ratio to crustal abundance of 21 in the bottom-sediment samples. The highest cadmium concentration (19 ppm) occurred in the same sample as the highest zinc concentration (GLOC-8, confluence of Spring and Neosho Rivers).

The average arsenic ratio to crustal abundance was 4.5, although absolute concentrations were generally less than 10 ppm.

The lead concentrations in the bottom-sediment samples are moderate, averaging only 43 ppm. Although these samples were collected relatively close to the former mining districts to the north, the lead concentrations are only three times crustal abundance (13 ppm) in the sediment. One of the two primary commodities mined was lead. To that end, producers tried to retain as much lead as possible in the milled and refined product and not release notable quantities of lead in the tailings. Second, the specific gravity of galena, the primary lead-bearing mineral, is high at about 7.5, so any galena released in the mining or milling stages likely would remain close to the initial source in the stream system as detrital grains. In addition, galena is not very soluble under the near-neutral conditions common in these stream waters (Weast, 1978). Conversely, zinc is highly mobile in the aqueous phase once it has been dissolved from sphalerite (Stumm and Morgan, 1981). Thus, zinc has the ability to move relatively far and quickly and can settle into bed sediment sorbed to colloidal iron phases or coprecipitated with other phases.

The total carbon content in the bottom-sediment samples ranged from 0.86 percent to 4.6 percent. The carbonate carbon content was low, averaging 0.06 percent, and ranging from 0.01 percent to 0.23 percent, with a median concentration of 0.03 percent. Thus, carbon was mostly in the organic form within the sediments.

Core GLOC-9 Elk River Arm of Grand Lake O' the Cherokees (Table 1-3)

The core site in the Elk River arm of Grand Lake O' the Cherokees was chosen because the Elk River drains a watershed that does not contain Tri-State mining districts, and the site was expected to define local background concentrations for mining-related elements. This core seemed to have minimal, if any, influence from mining-related elements.

The estimated SiO₂ content in core GLOC-9 averaged 88 percent. All of the other major element concentrations were less than crustal abundance. Calcium and magnesium concentration ratios were at 8 and 9 percent of crustal abundance, respectively. The median iron concentration ratio was 27 percent, whereas the ratio for sodium was 10 percent of crustal abundance. Aluminum and potassium median concentrations were greater, each at about 50 percent of crustal abundance, probably reflecting the clay content of the sediment.

Trace-element concentrations in core GLOC-9 were uniformly small, although arsenic (median 4 ppm) and cadmium (median 0.3 ppm) concentrations were two times crustal abundance. Lead concentrations averaged slightly greater than crustal abundance at 16 ppm. The copper concentrations from the background core averaged 12 ppm, less than the crustal abundance concentration of 68 ppm.

There were no clear trends with depth for As, Cd, or Pb in this core. Figure 1–1 shows the concentration versus depth profile for Cd, Pb, and Zn for core GLOC-9.

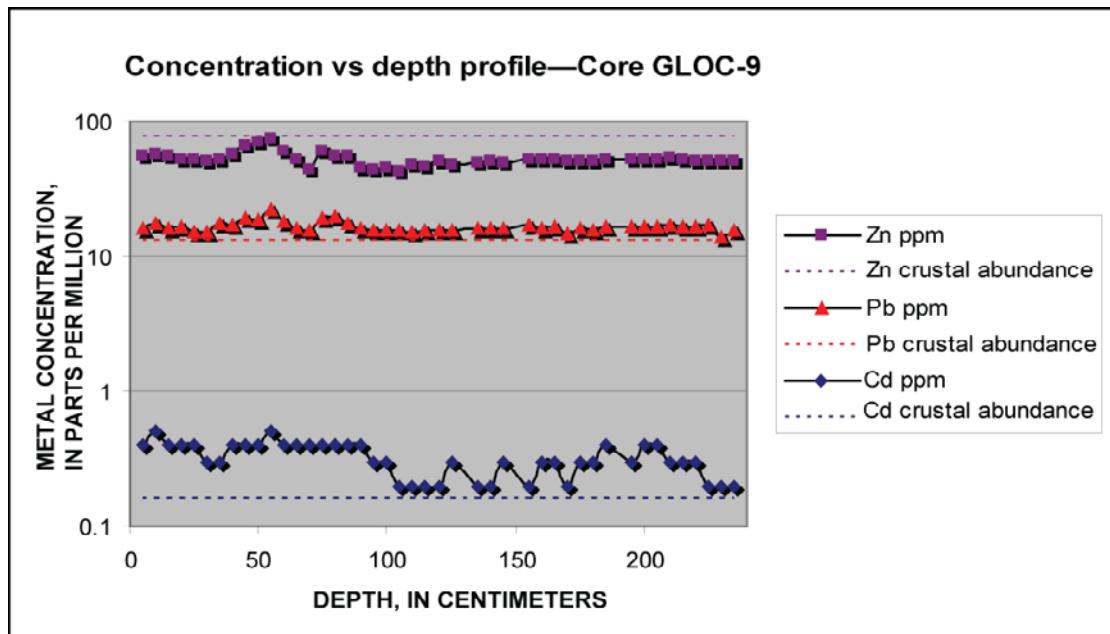


Figure 1–1. Concentration versus depth profile for cadmium (Cd), lead (Pb), and zinc (Zn) for background core GLOC-9.

Zinc concentrations in this core averaged 52 ppm, below crustal abundance, but there was some variability in the upper part of the core. The top three intervals of this core had an average zinc concentration of 52 ppm. At a depth of 55 cm (22 in), there was a maximum concentration of 74 ppm, which was followed by an irregular decrease of concentrations to 45 ppm at 90-cm (35-in) depth. From there to the bottom of the core (235 cm [93 in]), the zinc concentration averaged 50 ppm.

Core GLOC-2 Neosho River above Reservoir (Table 1–4)

Core GLOC-2 was collected on the Neosho River about 1 km (0.6 mi) upstream from its confluence with the Spring River. Although the surface samples from this core, discussed previously, contained near-background concentrations of most elements, subsamples deeper in the core contained higher concentrations of As, Cd, and Zn. Figure 1–2 shows the concentration versus depth profile for Cd, Pb, and Zn for core GLOC-2.

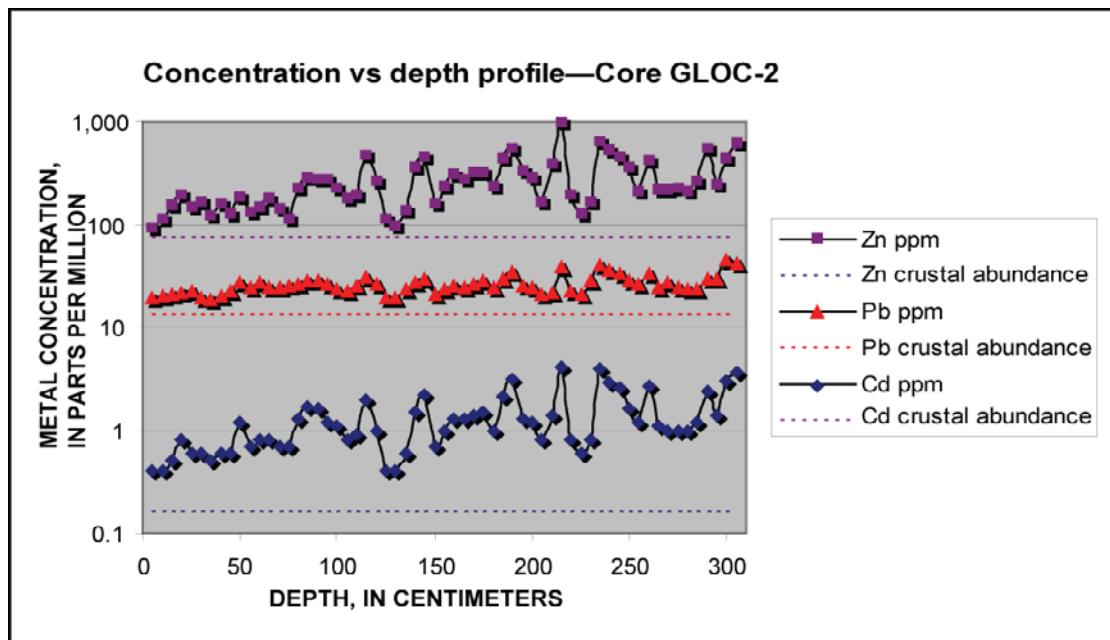


Figure 1–2. Concentration versus depth profile for cadmium (Cd), lead (Pb), and zinc (Zn) for core GLOC-2.

Arsenic concentrations ranged from 4 to 15 ppm, with the median concentration (7 ppm) having a ratio to crustal abundance of 3.9. The cadmium concentrations ranged from 0.4 to 4.1 ppm, with the median concentration (1.1 ppm) resulting in a ratio to crustal abundance of 7.3 and a ratio to the local background determined from core GLOC-9 of 3.6. Lead concentrations were only slightly greater than the background concentration, with the high concentration being 45 ppm at a depth of 300 cm (118 in). The ratio of the median lead concentration (25 ppm) to the background concentration was 1.9, about half that for the surface sediment samples.

The uppermost interval from core GLOC-2 had a zinc concentration of 95 ppm. Down to a depth of 65 cm (26 in), the zinc concentration averaged 160 ppm. With increasing depth, zinc concentrations increased irregularly, reaching a peak of 1,000 ppm at a depth of 215 cm (85 in). The median zinc concentration for the whole core was 230 ppm—three times crustal abundance and 4.4 times the local background as determined from core GLOC-9.

The major element concentrations and ratios to crustal abundance in this core were similar to those of the surface sediment samples discussed previously.

Core GLOC-3 Spring River above Reservoir (Table 1-5)

Core GLOC-3 was collected on the Spring River about 1.6 km (1 mi) upstream from the confluence with the Neosho River. This core had numerous subsamples with concentrations of Cd, Pb, and Zn far greater than crustal abundance levels and had definitive trends with depth. The major element concentration ratios to crustal abundance of this core were similar to those of the surface sediments.

Figure 1–3 shows the concentration versus depth profile for Cd, Pb, and Zn for core GLOC-3. Cadmium concentrations in the top 50 cm (20 in) were about 10 ppm and then increased to over 30 ppm between 80- and 90-cm depth (31–35 in); this value is about 200 times the crustal abundance of 0.16 ppm. Cadmium concentrations declined irregularly with depth in the core to 0.6 ppm at 145 cm (57 in). Below 145 cm, the concentrations remained steady at 0.3 ppm, or only twice crustal abundance. It is likely that core GLOC-3 penetrated to material deposited before the deposition of mining-related sediments in the Spring River.

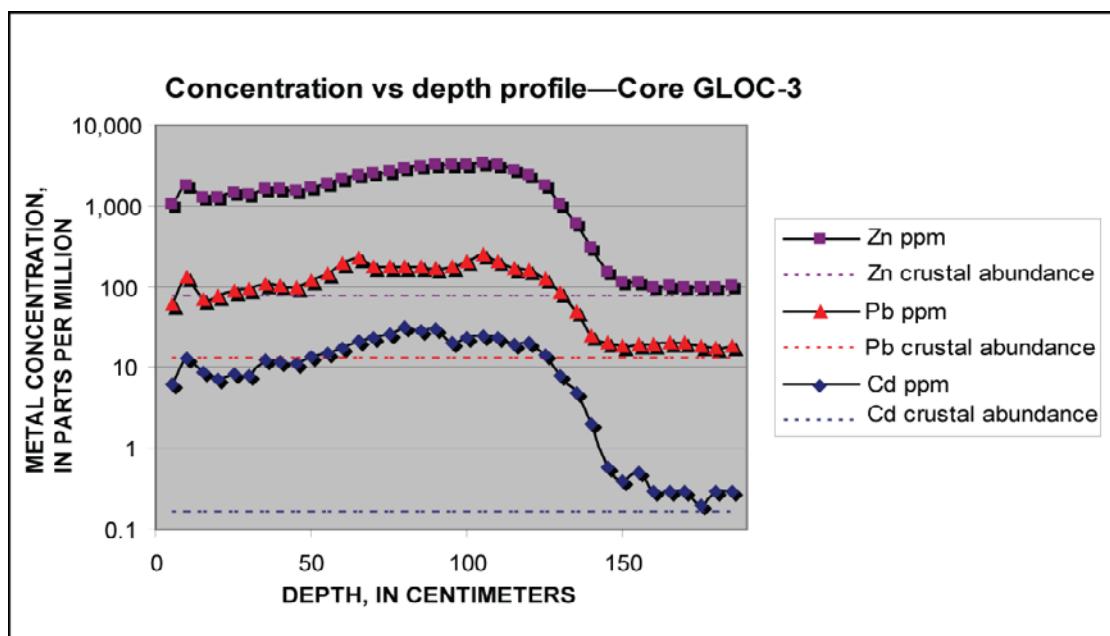


Figure 1–3. Concentration versus depth profile for cadmium (Cd), lead (Pb), and zinc (Zn) for core GLOC-3.

The depth versus concentration profile for lead was similar to that of cadmium. The top 50 cm (20 in) of sediment had an average lead concentration of about 95 ppm, about 6 times crustal abundance, and more than twice the average ratio for all surface sediment. At a depth of 65 cm (26 in), the lead concentration was 226 ppm, after which it decreased to about 170 ppm until a depth of 94 cm (37 in). The lead concentration increased to 250 ppm at a depth of 105 cm (41 in), and then decreased to 20 ppm, near crustal abundance, from 145 cm (57 in) to the bottom of the core. The lead data from this core indicate that most of the lead entering the Grand Lake O' the Cherokees is coming from the Spring River.

Similarly, zinc concentrations in this core are greater than those of any other sediment samples. The topmost interval of this core had a zinc concentration of 1,020 ppm; the concentrations then increased to 3,340 ppm at a depth of 105 cm (41 in). Zinc concentrations then decreased to about 100

ppm from a depth of 145 cm (57 in) down to the bottom of the core (185 cm [73 in]). The bottom sample concentrations were only 1.4 times crustal abundance, indicating that this core penetrated to below sediments affected by mining. The larger lead and zinc concentrations in this core, relative to those of the core from the Neosho River, GLOC-2, may be related to the fact that a larger percentage of the land area of the Spring River basin had been mined compared to the percentage of area mined in the Neosho River basin.

Core GLOC-7 Grand Lake O' the Cherokees Reservoir (Table 1-6)

Core GLOC-7 was collected from a site about 3.5 km (2.2 mi) downstream from the confluence of the Neosho and Spring Rivers, in the main body of the reservoir. The purpose of collecting this core was to document metal depth versus concentration profiles in Grand Lake O' the Cherokees. Both cadmium and zinc showed distinctive profiles. The major element concentration ratios to crustal abundance in this core were similar to those of the surface sediments.

Figure 1-4 shows the concentration versus depth profile for Cd, Pb, and Zn for core GLOC-7. Whereas the cadmium concentrations were much less than those from core GLOC-3, they had a distinct bimodal pattern. Between the top of the core and a depth of 70 cm (28 in), the cadmium concentrations oscillate between 1 and 4 ppm. From 70 cm to the bottom of the core at 135 cm (53 in), the cadmium concentrations decrease to about 0.4 ppm, only 2.6 times crustal abundance. This core penetrated to below mining-affected sediment, possibly into pre-reservoir soil that was flooded and covered with sediment after the completion of the Pensacola Dam.

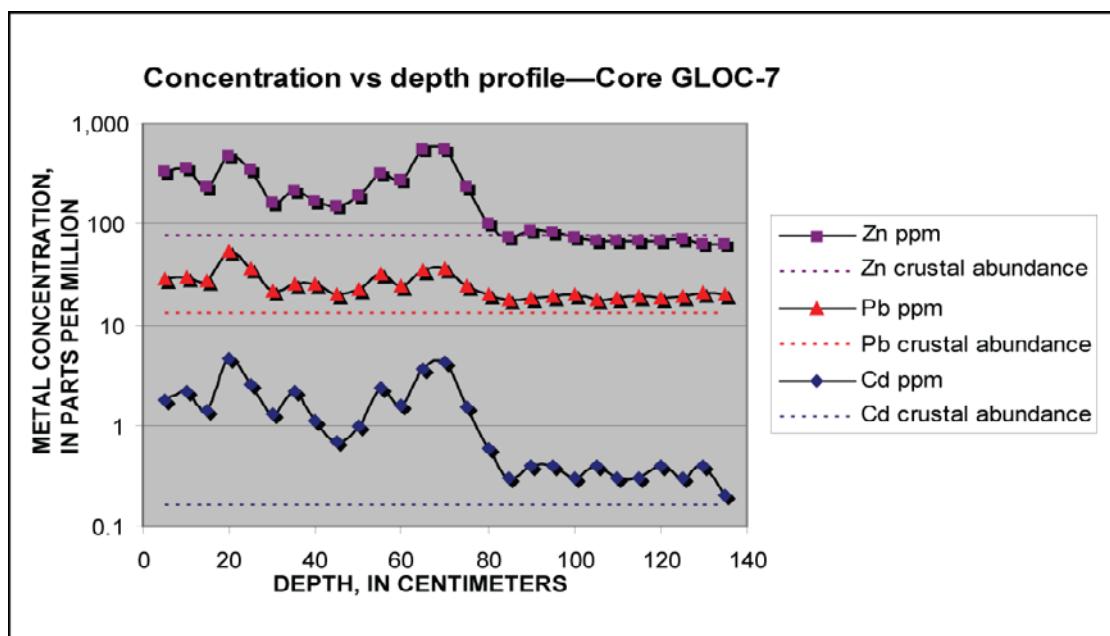


Figure 1-4. Concentration versus depth profile for cadmium (Cd), lead (Pb), and zinc (Zn) for core GLOC-7.

Lead concentrations in core GLOC-7 are not highly elevated above background. Concentrations are irregular between the top interval (28 ppm), ranging up to 54 ppm at 20-cm (8-in) depth and decreasing to 20 ppm at a depth of 80 cm (31 in), which is 1.5 times crustal abundance. Despite large concentrations entering the reservoir from the Spring River, the lead concentrations are attenuated by the time sediment reaches GLOC-7, probably due to the immobility of lead.

Zinc concentrations in core GLOC-7 also had a bimodal distribution, with one depth range of elevated and variable concentrations and a second depth range with values near crustal abundance. Beneath the top sample interval, which has a zinc concentration of 336 ppm (crustal abundance is 76 ppm), the zinc concentration profile ranges from a local maximum of 463 ppm at 20-cm (8-in) depth, down to a local minimum of 152 ppm at 45 cm (18 in), and then up to a core maximum of 554 ppm at 70-cm (28-in) depth, after which the concentrations show a sharp decline. Below 85-cm (33-in) depth, zinc concentration remained about 70 ppm, the crustal abundance concentration. The zinc concentration profile, then, also appears to indicate that the core penetrated to pre-reservoir material.

Core GLOC-8 Confluence of Neosho and Spring Rivers (Table 1-7)

Core GLOC-8 was collected just off the tip of land that protrudes into the reservoir on the southern end of Twin Bridges State Park. The site was selected to represent a possible mixing zone for sediment from the Neosho and Spring Rivers; however, there may not have been equal contributions of sediment from each river at this site. Metals concentrations of the top three intervals indicate that the Spring River contributes more to the current surface sediment signature (see the previous discussion on bottom-surface sediment). The major element concentration ratios to crustal abundance in this core were similar to those of the surface sediments.

Figure 1–5 shows the concentration versus depth profile for Cd, Pb, and Zn for core GLOC-8. The cadmium concentrations at the top three intervals (15 cm [6 in]) of this core are about 20 ppm, 130 times crustal abundance. At 25-cm (10-in) depth, a maximum of 25 ppm cadmium was measured, followed by a decrease to 3.1 ppm at 50-cm (20-in) depth. From 50 cm to the bottom of the core (75 cm [30 in]), cadmium concentrations decreased to 0.7 ppm, or about 4.5 times crustal abundance. The background core GLOC-9 had cadmium concentrations of about 0.3 ppm, so it appears that core GLOC-8 cadmium concentrations were approaching local background concentrations near the base.

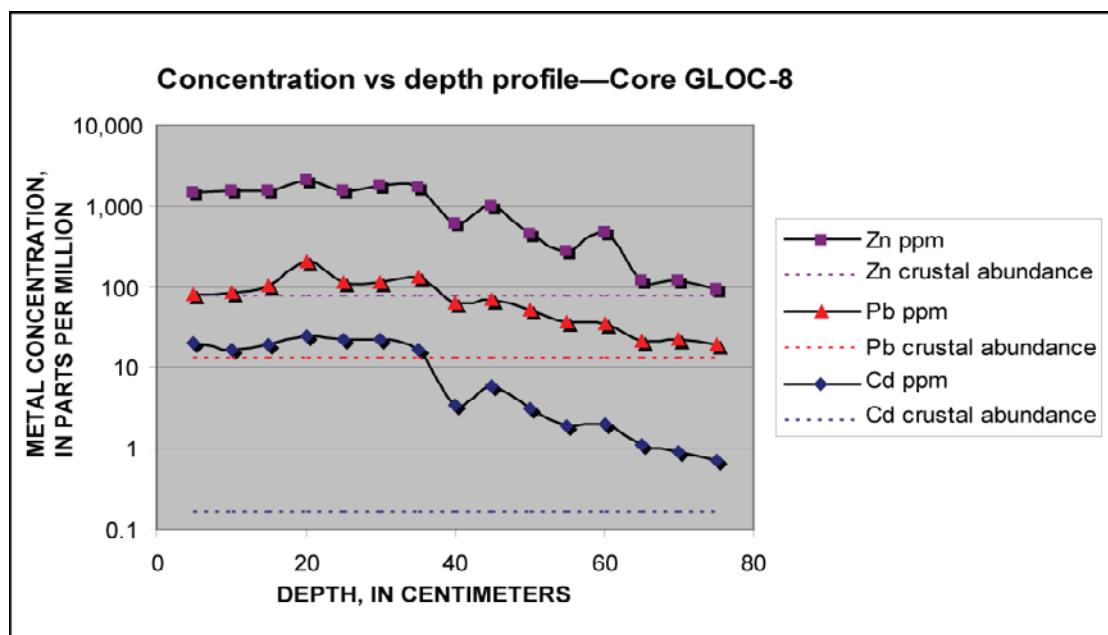


Figure 1-5. Concentration versus depth profile for cadmium (Cd), lead (Pb), and zinc (Zn) for core GLOC-8.

Lead concentrations in core GLOC-8 were also elevated, relative to background. This site is only 1.6 km (1 mi) downstream from GLOC-3, the Spring River core. The first three intervals of this core averaged 90 ppm lead, followed by a maximum of 204 ppm at a depth of 20 cm (8 in). The maximum lead concentration from the Spring River core (GLOC-3) was 250 ppm. From 20-cm depth to the core bottom at 75 cm (30 in), there is an irregular decrease in lead concentrations, with the bottom three intervals containing 20 ppm, or just 1.4 times crustal abundance.

Zinc concentrations in the upper 60 cm (24 in) of core GLOC-8 were much greater than background concentrations. The top three intervals of this core contained about 1,500 ppm zinc, with a concentration of 2,030 ppm occurring at a depth of 20 cm (8 in). The relatively large zinc concentrations in this core appear to be contributed mostly by sediment from the Spring River, since GLOC-2, the core from the Neosho River, contained smaller zinc concentrations than core GLOC-8. Between depths of 25 and 35 cm (10–14 in), the zinc concentration in core GLOC-8 averaged 1,690 ppm and then decreased irregularly to 478 ppm at a depth of 60 cm (24 in). In the last three depth intervals, 65 to 75 cm (26–30 in), the zinc concentration in this core averaged 110 ppm, which may be a result of the core's having penetrated to below mining-affected sediment.

Summary and Conclusion

Grand Lake O' the Cherokees is a reservoir downstream from the Tri-State District of Oklahoma, Kansas, and Missouri. This is a Mississippi Valley-type deposit, formerly one of the largest producers of lead and zinc in the world. Sediment core samples and bottom sediment samples were collected from the reservoir and from the Neosho River and the Spring River and analyzed to assess their geochemical signatures.

The Elk River arm (core GLOC-9) to the reservoir defines local background concentrations for the metals Zn, Pb, Cd, As, and Cu. The background for zinc was 52 ppm, which is less than the crustal abundance of 76 ppm. The local background for lead was 16 ppm, similar to the crustal abundance concentration of 13 ppm. The local background concentration for cadmium was 0.3 ppm, about twice that of the crustal abundance of 0.16 ppm. The local background concentration for arsenic was 4.2 ppm, compared to 1.8 ppm crustal abundance. Copper had a background concentration of 12 ppm, compared to 68 ppm crustal abundance.

The sediment samples collected from the sediment-water interface had larger average concentrations of Zn, Cd, and Pb than local background. The zinc concentrations of the bottom-sediment samples averaged 500 ppm, or almost 10 times the local background. The bottom-sediment lead concentrations averaged 42 ppm, about 2.6 times the local background. The cadmium concentrations of the bottom-sediment samples averaged 2.5 ppm, about 8 times local background. Locally, terrestrial sediment input into the upper parts of some coves can dilute the sediment signature.

The Spring River core GLOC-3 had the largest concentrations of mining-related elements. These concentrations were larger than those from the Neosho River core GLOC-2, possibly related to the fact that a larger percentage of the land area in the Spring River basin was mined compared to the percentage mined in the Neosho River Basin. The core collected from just south of Twin Bridges State Park, core GLOC-8 at the confluence of the Spring and Neosho Rivers, showed a mixing zone with more mining-related elements coming from the Spring River side.

The element zinc showed the most definitive patterns in concentration versus depth profiles. The core collected from the main body of the reservoir, GLOC-7, showed affected sediment down to a depth of 85 cm (33 in), below which mining-related element concentrations approached local background levels. This core and cores GLOC-3 and GLOC-8 appear to have penetrated to below mining-affected sediment.

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Appendix 1. Analytical Data

Table 1–1. Elements analyzed, instrument used, reporting limits, and mean of eight procedure blanks for four-acid digestion procedure. ([Click to see Excel file.](#))

[ICP-AES, inductively coupled plasma–atomic emission spectroscopy; ICP-MS, inductively coupled plasma–mass spectroscopy; %, percent; ppm, parts per million; n/a, not available]

Element	Instrument	Unit	Reporting limit	Blank mean (n=8)
Carb C	Coulometric	percent	0.003	<0.003
Total C	Combustion	percent	0.01	n/a
Al	ICP-AES	percent	0.01	<0.01
Ca	ICP-AES	percent	0.01	<0.01
Fe	ICP-AES	percent	0.01	<0.01
K	ICP-AES	percent	0.01	<0.01
Mg	ICP-AES	percent	0.01	<0.01
Na	ICP-AES	percent	0.01	<0.01
S	ICP-AES	percent	0.01	<0.01
Ti	ICP-AES	percent	0.01	<0.01
As	ICP-MS	ppm	1.0	< 1
Ba	ICP-AES	ppm	5.0	< 5
Bi	ICP-MS	ppm	0.04	< 0.04
Cd	ICP-MS	ppm	0.1	< 0.1
Co	ICP-MS	ppm	0.1	0.2
Cr	ICP-AES	ppm	1.0	2.0
Cu	ICP-AES	ppm	0.5	0.65
Ga	ICP-MS	ppm	0.05	0.06
In	ICP-MS	ppm	0.02	< 0.02
Li	ICP-AES	ppm	1.0	< 1
Mn	ICP-AES	ppm	5.0	< 5
Ni	ICP-AES	ppm	0.5	0.60
P	ICP-AES	ppm	50	< 50
Pb	ICP-MS	ppm	0.5	0.80
Rb	ICP-MS	ppm	0.2	0.30
Sr	ICP-AES	ppm	0.5	< 0.5
V	ICP-AES	ppm	1.0	< 1
Zn	ICP-AES	ppm	1.0	< 1

Table 1–2. Grand Lake O' the Cherokees bottom-sediment sample analyses. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon. First five samples are composites of top three intervals of cores GLOC-2, GLOC-3, GLOC-7, GLOC-8, and GLOC-9. ([Click to see Excel file.](#))

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; %, percent; ppm, parts per million]

Lab no.	Field no.	Total C %	Carbonate C %	Organic C %	Al %	Ca %	Fe %	K %
composite	GLOC-2_1-7	1.01	0.04	0.97	5.35	0.62	2.26	1.54
composite	GLOC-3_1-7	1.39	0.08	1.30	2.87	0.48	1.65	0.71
composite	GLOC-7_1-7	1.68	0.04	1.65	6.60	0.85	2.95	1.43
composite	GLOC-8_1-7	1.13	0.01	1.11	5.20	0.473	2.35	1.32
composite	GLOC-9_1-7	1.54	0.24	1.30	2.82	1.02	1.16	0.99
C-310716	GLOC-12_1-7	1.79	0.02	1.77	7.39	0.83	3.46	1.67
C-310690	GLOC-13_1-7	1.25	0.01	1.24	4.81	0.62	2.08	1.10
C-310708	GLOC-14_1-7	2.01	0.08	1.93	7.60	1.09	3.3	1.55
C-310685	GLOC-15_1-7	1.99	0.02	1.97	8.11	0.94	3.91	1.76
C-310703	GLOC-16_1-7	1.98	0.19	1.79	4.89	1.23	2.22	1.27
C-310681	GLOC-17_1-7	1.77	0.04	1.73	7.41	0.94	3.51	1.69
C-310713	GLOC-18_1-7	1.29	0.03	1.26	5.69	0.65	2.39	1.59
C-310707	GLOC-19_1-7	1.76	0.03	1.73	7.71	0.89	3.57	1.70
C-310688	GLOC-20_1-7	1.78	0.01	1.77	3.33	0.41	1.64	0.90
C-310702	GLOC-21_1-7	2.05	0.10	1.95	8.32	1.26	3.92	1.77
C-310684	GLOC-22_1-7	1.57	0.02	1.55	7.58	0.83	3.45	1.77
C-310700	GLOC-23_1-7	0.86	0.01	0.85	4.37	0.32	1.81	1.15
C-310693	GLOC-24_1-7	1.36	0.06	1.30	4.68	0.71	2.15	1.31
C-310712	GLOC-25_1-7	1.36	0.02	1.34	5.93	0.61	2.57	1.57
C-310710	GLOC-26_1-7	4.58	0.23	4.35	6.14	1.74	3.12	1.47

Table 1–2. Grand Lake O' the Cherokees bottom-sediment sample analyses. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon. First five samples are composites of top three intervals of cores GLOC-2, GLOC-3, GLOC-7, GLOC-8, and GLOC-9.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; %, percent; ppm, parts per million]

Lab no.	Field no.	Mg %	Na %	S %	Ti %	As ppm	Ba ppm	Bi ppm
composite	GLOC-2_1-7	0.45	0.64	0.03	0.23	6	500	0.23
composite	GLOC-3_1-7	0.17	0.23	0.07	0.15	4	269	0.16
composite	GLOC-7_1-7	0.35	0.65	0.05	0.21	6	450	0.19
composite	GLOC-8_1-7	0.38	0.46	0.04	0.24	7	465	0.27
composite	GLOC-9_1-7	0.19	0.22	0.04	0.17	3	297	0.14
C-310716	GLOC-12_1-7	0.63	0.33	0.12	0.31	10	547	0.33
C-310690	GLOC-13_1-7	0.34	0.30	0.16	0.23	7	414	0.28
C-310708	GLOC-14_1-7	0.62	0.24	0.19	0.29	10	532	0.35
C-310685	GLOC-15_1-7	0.71	0.32	0.11	0.30	10	604	0.36
C-310703	GLOC-16_1-7	0.36	0.28	0.14	0.22	8	434	0.22
C-310681	GLOC-17_1-7	0.64	0.38	0.12	0.25	10	574	0.34
C-310713	GLOC-18_1-7	0.46	0.60	0.07	0.26	7	500	0.25
C-310707	GLOC-19_1-7	0.66	0.35	0.13	0.30	11	566	0.33
C-310688	GLOC-20_1-7	0.23	0.24	0.09	0.19	6	338	0.18
C-310702	GLOC-21_1-7	0.74	0.27	0.17	0.28	13	577	0.36
C-310684	GLOC-22_1-7	0.67	0.42	0.11	0.28	9	583	0.34
C-310700	GLOC-23_1-7	0.25	0.41	0.03	0.24	5	487	0.20
C-310693	GLOC-24_1-7	0.36	0.32	0.08	0.23	7	447	0.23
C-310712	GLOC-25_1-7	0.49	0.53	0.06	0.24	7	518	0.27
C-310710	GLOC-26_1-7	0.56	0.32	0.33	0.24	16	511	0.36

Table 1–2. Grand Lake O' the Cherokees bottom-sediment sample analyses. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon. First five samples are composites of top three intervals of cores GLOC-2, GLOC-3, GLOC-7, GLOC-8, and GLOC-9.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; %, percent; ppm, parts per million]

Lab no.	Field no.	Cd ppm	Co ppm	Cr ppm	Cu ppm	Ga ppm	In ppm	Li ppm	Mn ppm
composite	GLOC-2_1-7	0.43	11	51	15.0	13	0.04	30	488
composite	GLOC-3_1-7	9.2	9.3	34	16.5	6.93	0.04	20	543
composite	GLOC-7_1-7	1.8	8.9	40	16.6	10.5	0.04	25	378
composite	GLOC-8_1-7	19	11.3	50	23.0	11.9	0.07	31	534
composite	GLOC-9_1-7	0.40	7.7	40	11.5	6.0	0.02	17	307
C-310716	GLOC-12_1-7	3.1	13.9	67	25.5	17	0.06	48	647
C-310690	GLOC-13_1-7	1.7	9.5	57	19.5	10.5	0.04	31	346
C-310708	GLOC-14_1-7	2.3	14.0	77	24.5	17.6	0.07	49	382
C-310685	GLOC-15_1-7	2.7	14.2	74	26.7	18.4	0.07	52	1,040
C-310703	GLOC-16_1-7	2.2	11.9	54	19.9	11.2	0.04	31	510
C-310681	GLOC-17_1-7	3.4	14.5	77	27.1	17.3	0.07	47	859
C-310713	GLOC-18_1-7	2.8	10.5	48	18.2	12.1	0.05	34	503
C-310707	GLOC-19_1-7	3.4	14.4	69	25.1	18.0	0.07	50	735
C-310688	GLOC-20_1-7	0.70	8.1	38	13.8	7.3	0.03	22	281
C-310702	GLOC-21_1-7	2.8	16.3	137	30.0	20.1	0.08	54	690
C-310684	GLOC-22_1-7	3.0	13.8	65	25.9	17.0	0.07	48	603
C-310700	GLOC-23_1-7	1.0	11.2	39	14.1	9.9	0.04	26	559
C-310693	GLOC-24_1-7	2.7	12.7	38	18.9	10.8	0.04	30	503
C-310712	GLOC-25_1-7	2.7	11.1	53	22.2	13.0	0.05	35	514
C-310710	GLOC-26_1-7	3.8	22.2	63	30.0	14.1	0.06	40	795

Table 1–2. Grand Lake O' the Cherokees bottom-sediment sample analyses. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon. First five samples are composites of top three intervals of cores GLOC-2, GLOC-3, GLOC-7, GLOC-8, and GLOC-9.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; %, percent; ppm, parts per million]

Lab no.	Field no.	Ni ppm	P ppm	Pb ppm	Rb ppm	Sr ppm	V ppm	Zn ppm
composite	GLOC-2_1-7	22.0	607	20.2	90.0	113	78	122
composite	GLOC-3_1-7	16.4	733	87.0	46.8	42.0	45	1,370
composite	GLOC-7_1-7	18.6	530	28.3	77.3	92.7	65	310
composite	GLOC-8_1-7	24.7	747	90.0	82.0	75.0	79	1,523
composite	GLOC-9_1-7	22.2	357	17.0	49.0	39.3	42	56
C-310716	GLOC-12_1-7	33.0	970	46.9	117	103	119	574
C-310690	GLOC-13_1-7	28.1	560	34.4	71.5	69.4	73	402
C-310708	GLOC-14_1-7	34.8	840	47.0	119	94.5	120	540
C-310685	GLOC-15_1-7	34.6	1,370	45.2	118	107	127	516
C-310703	GLOC-16_1-7	26.1	650	37.0	80.1	78.7	77	448
C-310681	GLOC-17_1-7	34.2	1,020	48.9	111	106	115	562
C-310713	GLOC-18_1-7	23.4	680	36.2	87.7	100	82	450
C-310707	GLOC-19_1-7	33.2	1,150	46.4	123	109	118	577
C-310688	GLOC-20_1-7	17.4	600	22.4	58.9	51.5	57	242
C-310702	GLOC-21_1-7	72.2	1,170	46.2	132	114	135	550
C-310684	GLOC-22_1-7	32.1	950	42.1	112	113	116	528
C-310700	GLOC-23_1-7	19.9	360	21.2	75.3	68.3	62	134
C-310693	GLOC-24_1-7	22.0	670	37.1	76.0	78.1	73	478
C-310712	GLOC-25_1-7	25.1	790	36.9	92.5	97.4	85	473
C-310710	GLOC-26_1-7	40.2	1,360	76.2	97.9	108	122	1,020

Table 1–3. Core sediment background sample analyses for GLOC-9, Elk River arm of Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon. ([Click to see Excel file.](#))
 [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Depth in inches	Depth in cm	Total C %	Carbonate C %	Organic C %	Al %	Ca %	Fe %
C-310665	GLOC-9_1-3	2	5	1.6	0.31	1.29	2.71	1.23	1.13
C-310655	GLOC-9_3-5	4	10	1.53	0.23	1.3	2.94	0.99	1.19
C-310627	GLOC-9_5-7	6	15	1.49	0.19	1.3	2.82	0.84	1.15
C-310670	GLOC-9_7-9	8	20	1.47	0.2	1.27	2.67	0.88	1.04
C-310671	GLOC-9_9-11	10	25	1.35	0.19	1.16	2.47	0.83	1.02
C-310661	GLOC-9_11-13	12	30	1.42	0.18	1.24	2.7	0.78	1.09
C-310650	GLOC-9_13-15	14	35	1.41	0.11	1.3	2.66	0.58	1.11
C-310642	GLOC-9_15-17	16	40	1.6	0.1	1.5	2.87	0.6	1.14
C-310672	GLOC-9_17-19	18	45	1.98	0.1	1.88	3.14	0.65	1.24
C-310647	GLOC-9_19-21	20	50	3.36	0.06	3.3	3.27	0.55	1.31
C-310639	GLOC-9_21-23	22	55	2.04	0.22	1.82	3.46	0.98	1.43
C-310628	GLOC-9_23-25	24	60	1.53	0.1	1.43	3.05	0.58	1.28
C-310629	GLOC-9_25-27	26	65	1.16	0.08	1.08	2.46	0.4	1.16
C-310645	GLOC-9_27-29	28	70	0.68	<0.003	0.68	3.17	0.17	1.21
C-310668	GLOC-9_29-31	30	75	1.14	0.02	1.12	3.02	0.25	1.24
C-310653	GLOC-9_31-33	32	80	1.19	<0.003	1.19	3.19	0.22	1.24
C-310644	GLOC-9_33-35	34	85	1.03	0.05	0.98	2.73	0.32	1.18
C-310649	GLOC-9_35-37	36	90	0.74	0.01	0.73	3.21	0.17	1.23
C-310632	GLOC-9_37-39	38	95	0.41	<0.003	0.41	3.26	0.16	1.29
C-310669	GLOC-9_39-41	40	100	0.43	<0.003	0.43	3.45	0.16	1.32
C-310660	GLOC-9_41-43	42	105	0.4	0.02	0.38	3.3	0.16	1.37
C-310640	GLOC-9_43-45	44	110	0.41	0.02	0.39	3.58	0.18	1.52
C-310636	GLOC-9_45-47	46	115	0.58	0.05	0.53	3.67	0.18	1.65
C-310659	GLOC-9_47-49	48	120	0.42	0.04	0.38	3.64	0.18	1.65
C-310656	GLOC-9_49-51	50	125	0.39	0.03	0.36	3.72	0.19	1.65
C-310630	GLOC-9_51-53	54	135	0.37	0.02	0.35	3.89	0.18	1.71
C-310654	GLOC-9_53-55	56	140	0.34	0.02	0.32	3.7	0.17	1.6
C-310651	GLOC-9_55-57	58	145	0.34	0.01	0.33	3.79	0.18	1.61
C-310641	GLOC-9_57-59	62	155	0.33	0.01	0.32	3.84	0.19	1.63
C-310658	GLOC-9_59-61	64	160	0.32	0.01	0.31	3.8	0.18	1.59
C-310662	GLOC-9_61-63	66	165	0.31	0.01	0.3	3.81	0.19	1.63
C-310643	GLOC-9_63-65	68	170	0.29	0.01	0.28	3.72	0.19	1.61
C-310675	GLOC-9_65-67	70	175	0.29	0.01	0.28	3.76	0.19	1.58
C-310631	GLOC-9_67-69	72	180	0.28	<0.003	0.28	3.79	0.19	1.57
C-310652	GLOC-9_69-71	74	185	0.28	<0.003	0.28	3.85	0.2	1.58
C-310633	GLOC-9_71-73	78	195	0.25	0.01	0.24	3.94	0.21	1.61
C-310637	GLOC-9_73-75	80	200	0.25	<0.003	0.25	3.87	0.2	1.58
C-310638	GLOC-9_75-77	82	205	0.24	<0.003	0.24	3.94	0.21	1.58
C-310666	GLOC-9_77-79	84	210	0.24	0.01	0.23	3.89	0.21	1.57
C-310635	GLOC-9_79-81	86	215	0.23	<0.003	0.23	3.88	0.21	1.56
C-310674	GLOC-9_81-83	88	220	0.23	<0.003	0.23	3.74	0.21	1.52
C-310667	GLOC-9_83-85	90	225	0.21	<0.003	0.21	3.77	0.21	1.54
C-310657	GLOC-9_85-87	92	230	0.22	0.01	0.21	3.72	0.21	1.54
C-310664	GLOC-9_87-89	94	235	0.21	<0.003	0.21	3.75	0.21	1.53

Table 1–3. Core sediment background sample analyses for GLOC-9, Elk River arm of Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	K %	Mg %	Na %	S %	Ti %	As ppm	Ba ppm	Bi ppm
C-310665	GLOC-9_1-3	0.95	0.19	0.21	0.05	0.16	3	284	0.13
C-310655	GLOC-9_3-5	1.03	0.2	0.23	0.04	0.18	3	311	0.14
C-310627	GLOC-9_5-7	0.98	0.18	0.22	0.04	0.17	3	295	0.14
C-310670	GLOC-9_7-9	0.94	0.17	0.21	0.04	0.16	3	283	0.13
C-310671	GLOC-9_9-11	0.89	0.15	0.21	0.03	0.16	3	270	0.11
C-310661	GLOC-9_11-13	0.98	0.16	0.23	0.03	0.17	3	292	0.11
C-310650	GLOC-9_13-15	0.95	0.17	0.22	0.02	0.17	3	288	0.13
C-310642	GLOC-9_15-17	0.95	0.18	0.23	0.03	0.18	3	301	0.12
C-310672	GLOC-9_17-19	1.08	0.2	0.24	0.04	0.19	3	329	0.18
C-310647	GLOC-9_19-21	1.04	0.21	0.23	0.05	0.19	4	343	0.15
C-310639	GLOC-9_21-23	1.11	0.22	0.23	0.04	0.2	5	352	0.17
C-310628	GLOC-9_23-25	0.98	0.18	0.24	0.03	0.19	4	333	0.14
C-310629	GLOC-9_25-27	0.83	0.15	0.18	0.02	0.16	4	275	0.12
C-310645	GLOC-9_27-29	1.13	0.19	0.25	0.01	0.18	4	360	0.15
C-310668	GLOC-9_29-31	1.04	0.18	0.23	0.02	0.18	4	337	0.14
C-310653	GLOC-9_31-33	1.1	0.2	0.24	0.02	0.19	4	363	0.16
C-310644	GLOC-9_33-35	0.95	0.16	0.22	0.02	0.17	4	302	0.12
C-310649	GLOC-9_35-37	1.11	0.19	0.25	0.01	0.19	4	363	0.15
C-310632	GLOC-9_37-39	1.1	0.2	0.24	0.01	0.18	4	356	0.15
C-310669	GLOC-9_39-41	1.17	0.2	0.24	0.01	0.19	4	367	0.15
C-310660	GLOC-9_41-43	1.09	0.2	0.22	0.01	0.18	4	352	0.15
C-310640	GLOC-9_43-45	1.15	0.22	0.23	0.01	0.19	5	349	0.15
C-310636	GLOC-9_45-47	1.16	0.22	0.23	0.01	0.19	5	347	0.16
C-310659	GLOC-9_47-49	1.12	0.22	0.22	0.01	0.19	5	342	0.16
C-310656	GLOC-9_49-51	1.14	0.23	0.22	0.01	0.19	5	339	0.15
C-310630	GLOC-9_51-53	1.18	0.24	0.22	0.01	0.2	5	346	0.16
C-310654	GLOC-9_53-55	1.14	0.23	0.22	0.01	0.19	4	339	0.17
C-310651	GLOC-9_55-57	1.16	0.23	0.23	0.01	0.19	4	341	0.15
C-310641	GLOC-9_57-59	1.19	0.24	0.23	0.01	0.2	5	347	0.16
C-310658	GLOC-9_59-61	1.17	0.24	0.23	0.01	0.19	5	353	0.17
C-310662	GLOC-9_61-63	1.16	0.24	0.23	<0.01	0.19	5	362	0.17
C-310643	GLOC-9_63-65	1.16	0.23	0.23	<0.01	0.19	5	351	0.16
C-310675	GLOC-9_65-67	1.18	0.24	0.23	<0.01	0.19	4	356	0.17
C-310631	GLOC-9_67-69	1.18	0.24	0.24	<0.01	0.2	5	360	0.16
C-310652	GLOC-9_69-71	1.21	0.24	0.25	<0.01	0.2	4	373	0.16
C-310633	GLOC-9_71-73	1.25	0.25	0.26	0.01	0.2	5	393	0.18
C-310637	GLOC-9_73-75	1.23	0.25	0.26	<0.01	0.2	5	387	0.2
C-310638	GLOC-9_75-77	1.27	0.25	0.27	0.01	0.21	5	392	0.16
C-310666	GLOC-9_77-79	1.25	0.25	0.27	0.01	0.2	5	408	0.16
C-310635	GLOC-9_79-81	1.22	0.25	0.27	<0.01	0.19	5	391	0.16
C-310674	GLOC-9_81-83	1.21	0.24	0.27	<0.01	0.21	4	389	0.2
C-310667	GLOC-9_83-85	1.22	0.24	0.27	<0.01	0.2	5	390	0.17
C-310657	GLOC-9_85-87	1.19	0.24	0.26	<0.01	0.2	4	383	0.15
C-310664	GLOC-9_87-89	1.2	0.24	0.27	<0.01	0.21	4	385	0.16

Table 1–3. Core sediment background sample analyses for GLOC-9, Elk River arm of Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Cd ppm	Co ppm	Cr ppm	Cu ppm	Ga ppm	In ppm	Li ppm	Mn ppm
C-310665	GLOC-9_1-3	0.4	8	53	12.2	6	0.02	17	317
C-310655	GLOC-9_3-5	0.5	7.6	28	10.9	6	0.02	18	301
C-310627	GLOC-9_5-7	0.4	7.4	38	11.3	6	0.02	17	302
C-310670	GLOC-9_7-9	0.4	7.1	41	11.1	6	0.02	16	284
C-310671	GLOC-9_9-11	0.4	6.3	33	9.9	5.1	<0.02	16	287
C-310661	GLOC-9_11-13	0.3	6.5	29	10.5	5.3	<0.02	17	314
C-310650	GLOC-9_13-15	0.3	7.6	33	10.6	6	0.02	17	354
C-310642	GLOC-9_15-17	0.4	6.9	25	10.7	5.6	<0.02	18	355
C-310672	GLOC-9_17-19	0.4	7.8	27	12	6.9	0.03	20	428
C-310647	GLOC-9_19-21	0.4	7.2	45	12.3	6.9	0.03	20	474
C-310639	GLOC-9_21-23	0.5	9	36	13.7	7.9	0.03	21	802
C-310628	GLOC-9_23-25	0.4	7.6	33	11.7	6.3	0.02	19	580
C-310629	GLOC-9_25-27	0.4	7.2	24	10.1	5.4	0.02	15	683
C-310645	GLOC-9_27-29	0.4	8	52	12.3	7.4	0.03	19	378
C-310668	GLOC-9_29-31	0.4	9.9	34	12	6.8	0.03	18	388
C-310653	GLOC-9_31-33	0.4	10	30	13	7.1	0.03	19	414
C-310644	GLOC-9_33-35	0.4	9.3	30	11	6	0.02	18	599
C-310649	GLOC-9_35-37	0.4	8	38	12.2	7.3	0.03	19	369
C-310632	GLOC-9_37-39	0.3	8.4	36	11.3	7.4	0.03	19	360
C-310669	GLOC-9_39-41	0.3	8.2	35	11.4	7.4	0.03	20	385
C-310660	GLOC-9_41-43	0.2	9.8	65	11.9	8	0.03	20	570
C-310640	GLOC-9_43-45	0.2	8.4	35	11.3	8	0.03	21	695
C-310636	GLOC-9_45-47	0.2	9.3	37	11.9	8.1	0.03	21	1,110
C-310659	GLOC-9_47-49	0.2	9.8	32	11.6	8	0.03	21	1,060
C-310656	GLOC-9_49-51	0.3	9.9	37	12.3	7.9	0.03	21	980
C-310630	GLOC-9_51-53	0.2	11	35	12.4	8.7	0.03	22	798
C-310654	GLOC-9_53-55	0.2	10.5	37	12.2	8.6	0.03	21	754
C-310651	GLOC-9_55-57	0.3	9.4	32	12.8	8.1	0.03	21	745
C-310641	GLOC-9_57-59	0.2	8.9	36	12.8	8.5	0.03	22	704
C-310658	GLOC-9_59-61	0.3	8.7	44	12.4	8.9	0.03	22	727
C-310662	GLOC-9_61-63	0.3	9.3	43	12.8	9.1	0.03	21	738
C-310643	GLOC-9_63-65	0.2	9	35	12.6	7.9	0.03	21	799
C-310675	GLOC-9_65-67	0.3	9.6	32	12.5	8.4	0.03	21	798
C-310631	GLOC-9_67-69	0.3	9.7	40	12.9	8.4	0.03	22	897
C-310652	GLOC-9_69-71	0.4	9	38	12.6	8.5	0.03	21	869
C-310633	GLOC-9_71-73	0.3	10.2	68	14.1	9.1	0.04	22	892
C-310637	GLOC-9_73-75	0.4	9.7	36	13.6	8.9	0.03	22	875
C-310638	GLOC-9_75-77	0.4	9	39	13.7	8.6	0.03	22	818
C-310666	GLOC-9_77-79	0.3	10.1	36	13.9	8.9	0.03	22	911
C-310635	GLOC-9_79-81	0.3	9.2	49	13.2	8.6	0.03	22	880
C-310674	GLOC-9_81-83	0.3	9	34	17.6	8.5	0.03	21	824
C-310667	GLOC-9_83-85	0.2	9.8	68	13.3	8.6	0.03	21	857
C-310657	GLOC-9_85-87	0.2	8.1	46	12.4	7.4	0.03	21	825
C-310664	GLOC-9_87-89	0.2	9.4	35	12.8	8.4	0.03	21	809

Table 1–3. Core sediment background sample analyses for GLOC-9, Elk River arm of Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Ni ppm	P ppm	Pb ppm	Rb ppm	Sr ppm	V ppm	Zn ppm
C-310665	GLOC-9_1-3	30.8	340	16	46.8	39.2	41	55
C-310655	GLOC-9_3-5	16.2	360	17.7	49.3	40.2	42	57
C-310627	GLOC-9_5-7	19.7	370	16.1	51.1	38.4	43	55
C-310670	GLOC-9_7-9	20.4	320	16.5	46.9	36	40	53
C-310671	GLOC-9_9-11	16.6	290	15.1	43.4	35	35	52
C-310661	GLOC-9_11-13	15.3	320	15.2	46.6	37.8	38	51
C-310650	GLOC-9_13-15	17.5	320	17.4	47.4	35.8	40	53
C-310642	GLOC-9_15-17	15.8	330	17	48.7	37.7	41	57
C-310672	GLOC-9_17-19	16	410	19.1	54.9	40.4	46	66
C-310647	GLOC-9_19-21	18.6	460	18.8	59.3	39.6	50	71
C-310639	GLOC-9_21-23	19.5	510	22.1	59.3	42.2	54	74
C-310628	GLOC-9_23-25	18.1	430	18.2	53.5	39.4	46	61
C-310629	GLOC-9_25-27	14.1	350	16.2	42.6	32.3	37	52
C-310645	GLOC-9_27-29	25.9	310	15.8	61.2	40.3	51	44
C-310668	GLOC-9_29-31	19.4	360	19.3	54.8	38.2	47	60
C-310653	GLOC-9_31-33	20.3	360	19.9	59.5	40	49	55
C-310644	GLOC-9_33-35	20.1	320	17.3	47.6	36.6	40	55
C-310649	GLOC-9_35-37	21.4	320	16.1	60.1	40.5	52	45
C-310632	GLOC-9_37-39	20.1	290	15.6	59.4	38.7	51	44
C-310669	GLOC-9_39-41	19.2	270	15.5	60.8	40	54	45
C-310660	GLOC-9_41-43	33.7	270	15.5	62	37.5	53	43
C-310640	GLOC-9_43-45	20	270	15.2	63.4	38.1	56	48
C-310636	GLOC-9_45-47	20.1	270	15.6	63.1	38.2	56	46
C-310659	GLOC-9_47-49	20.1	250	15.4	64.7	36.6	58	50
C-310656	GLOC-9_49-51	22.6	260	15.8	60.1	37.4	58	48
C-310630	GLOC-9_51-53	21.7	260	16	65.4	38.9	60	49
C-310654	GLOC-9_53-55	21.7	250	16.2	64	36.9	60	50
C-310651	GLOC-9_55-57	19.7	250	15.9	62	38.1	57	49
C-310641	GLOC-9_57-59	21.6	240	16.8	64	39.1	59	53
C-310658	GLOC-9_59-61	23	220	16.2	66.1	38.6	60	52
C-310662	GLOC-9_61-63	23.8	230	16.5	65.8	38.7	61	52
C-310643	GLOC-9_63-65	24.2	210	14.8	64	38.6	60	51
C-310675	GLOC-9_65-67	20.5	210	16	62.8	38.5	57	51
C-310631	GLOC-9_67-69	21.8	230	15.8	62.5	40.9	59	51
C-310652	GLOC-9_69-71	21.9	230	16.5	63.9	41.2	59	52
C-310633	GLOC-9_71-73	35.5	240	16.3	67	43	62	53
C-310637	GLOC-9_73-75	22.6	220	16.3	64.6	42.3	60	53
C-310638	GLOC-9_75-77	23.8	220	16.5	65.2	43.7	60	52
C-310666	GLOC-9_77-79	26.3	220	17.1	66.3	43.9	61	54
C-310635	GLOC-9_79-81	23.8	210	16.5	63	43.4	58	52
C-310674	GLOC-9_81-83	21.9	210	16.6	65.5	42.4	60	51
C-310667	GLOC-9_83-85	36.8	220	16.8	62.6	43.3	60	50
C-310657	GLOC-9_85-87	23.3	210	14	60.8	42.6	57	51
C-310664	GLOC-9_87-89	20.6	210	15.8	62.5	43	58	50

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon. ([Click to see Excel file.](#))
 [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Depth in inches	Depth in cm	Total C %	Carbonate C %	Organic C %	Al %	Ca %	Fe %
C-312239	GLOC-2_1-3	2	5	1.01	0.07	0.94	5.42	0.70	2.34
C-312296	GLOC-2_3-5	4	10	1.01	0.03	0.98	5.28	0.59	2.20
C-312263	GLOC-2_5-7	6	15	1.02	0.03	0.99	5.35	0.57	2.24
C-312276	GLOC-2_7-9	8	20	1.03	0.03	1.00	5.24	0.55	2.25
C-312275	GLOC-2_9-11	10	25	1.15	0.03	1.12	5.46	0.57	2.33
C-312258	GLOC-2_11-13	12	30	0.98	0.02	0.96	5.04	0.48	2.06
C-312242	GLOC-2_13-15	14	35	0.79	0.02	0.77	4.65	0.46	1.79
C-312302	GLOC-2_15-17	16	40	0.96	0.03	0.93	4.96	0.53	2.17
C-312245	GLOC-2_17-19	18	45	0.91	0.02	0.89	5.42	0.49	2.25
C-312254	GLOC-2_19-21	20	50	1.08	0.02	1.06	5.87	0.54	2.54
C-312300	GLOC-2_21-23	22	55	0.96	0.01	0.95	5.25	0.48	2.34
C-312247	GLOC-2_23-25	24	60	0.89	0.02	0.87	5.36	0.45	2.36
C-312297	GLOC-2_25-27	26	65	0.93	0.01	0.92	5.05	0.44	2.22
C-312283	GLOC-2_27-29	28	70	0.91	0.02	0.89	5.44	0.50	2.37
C-312266	GLOC-2_29-31	30	75	0.78	0.01	0.77	5.01	0.47	2.04
C-312309	GLOC-2_31-33	32	80	0.98	0.02	0.96	5.06	0.49	2.18
C-312311	GLOC-2_33-35	34	85	1.07	0.02	1.05	5.58	0.53	2.42
C-312307	GLOC-2_35-37	36	90	1.08	0.01	1.07	5.36	0.51	2.46
C-312292	GLOC-2_37-39	38	95	1.63	0.03	1.60	4.95	0.52	2.11
C-312298	GLOC-2_39-41	40	100	1.1	0.08	1.02	4.88	0.62	2.31
C-312253	GLOC-2_41-43	42	105	0.92	0.05	0.87	5.29	0.54	2.18
C-312271	GLOC-2_43-45	44	110	1.25	0.01	1.24	5.94	0.65	2.67
C-312282	GLOC-2_45-47	46	115	1.21	0.03	1.18	5.49	0.56	2.50
C-312310	GLOC-2_47-49	48	120	1.11	0.03	1.08	5.73	0.55	2.37
C-312241	GLOC-2_49-51	50	125	0.88	0.1	0.78	5.18	0.63	2.11
C-312308	GLOC-2_51-53	52	130	0.83	0.07	0.76	4.94	0.58	2.01
C-312286	GLOC-2_53-55	54	135	1.13	0.07	1.06	5.76	0.65	2.52
C-312248	GLOC-2_55-57	56	140	1.04	0.13	0.91	5.9	0.54	2.51
C-312240	GLOC-2_57-59	58	145	1.24	0.01	1.23	6.18	0.54	2.71
C-312280	GLOC-2_59-61	60	150	0.74	0.07	0.67	4.56	0.54	1.88
C-312287	GLOC-2_61-63	62	155	0.75	0.06	0.69	4.67	0.53	1.97
C-312261	GLOC-2_63-65	64	160	0.98	0.06	0.92	5.29	0.60	2.39
C-312289	GLOC-2_65-67	66	165	0.92	0.03	0.89	4.95	0.46	2.06
C-312285	GLOC-2_67-69	68	170	1.05	0.04	1.01	5.34	0.53	2.24
C-312237	GLOC-2_69-71	70	175	1.2	0.04	1.16	6.08	0.61	2.64
C-312304	GLOC-2_71-73	72	180	1.15	0.05	1.10	5.67	0.62	2.63
C-312255	GLOC-2_73-75	74	185	1.42	0.11	1.31	6.27	0.75	2.83
C-312281	GLOC-2_75-77	76	190	1.37	0.08	1.29	6.06	0.71	2.68
C-312306	GLOC-2_77-79	78	195	1.43	0.12	1.31	6.22	0.87	3.06
C-312264	GLOC-2_79-81	80	200	1.19	0.13	1.06	5.59	0.79	2.47
C-312249	GLOC-2_81-83	82	205	0.79	0.03	0.76	4.73	0.65	1.93
C-312265	GLOC-2_83-85	84	210	1.1	0.13	0.97	5.46	0.81	2.29
C-312279	GLOC-2_85-87	86	215	1.41	0.17	1.24	5.97	0.95	2.71
C-312301	GLOC-2_87-89	88	220	1.16	0.2	0.96	5.69	1.1	2.38
C-312259	GLOC-2_89-91	90	225	1.15	0.16	0.99	5.58	0.99	2.47

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued
 [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Depth in inches	Depth in cm	Total C %	Carbonate C %	Organic C %	Al %	Ca %	Fe %
C-312290	GLOC-2_91-93	92	230	1.62	0.16	1.46	7.02	1.02	3.20
C-312238	GLOC-2_93-95	94	235	1.29	0.08	1.21	6.23	0.74	2.89
C-312303	GLOC-2_95-97	96	240	1.25	0.08	1.17	5.68	0.67	2.50
C-312251	GLOC-2_97-99	98	245	1.24	0.09	1.15	6.04	0.70	2.66
C-312236	GLOC-2_99-101	100	250	1.39	0.14	1.25	6.7	0.76	3.15
C-312293	GLOC-2_101-103	102	255	1.47	0.19	1.28	6.48	0.93	3.01
C-312277	GLOC-2_103-105	104	260	1.45	0.11	1.34	6.28	0.75	2.69
C-312262	GLOC-2_105-107	106	265	1.18	0.07	1.11	5.89	0.61	2.50
C-312246	GLOC-2_107-109	108	270	1.28	0.1	1.18	6.04	0.68	2.61
C-312291	GLOC-2_109-111	110	275	1.16	0.11	1.05	5.32	0.67	2.32
C-312256	GLOC-2_111-113	112	280	0.84	0.12	0.72	4.55	0.61	1.93
C-312299	GLOC-2_113-115	114	285	1.09	0.2	0.89	5.08	0.93	2.21
C-312295	GLOC-2_115-117	116	290	1.44	0.28	1.16	6.09	1.01	3.09
C-312274	GLOC-2_117-119	118	295	1.4	0.05	1.35	6.35	0.59	2.79
C-312250	GLOC-2_119-121	120	300	1.44	0.03	1.41	6.75	0.53	3.02
C-312272	GLOC-2_121-123	122	305	1.79	0.1	1.69	6.81	0.60	3.57

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	K %	Mg %	Na %	S %	Ti %	As ppm	Ba ppm
C-312239	GLOC-2_1-3	1.58	0.47	0.63	0.03	0.24	7	488
C-312296	GLOC-2_3-5	1.52	0.44	0.65	0.03	0.23	6	500
C-312263	GLOC-2_5-7	1.52	0.43	0.64	0.03	0.22	6	508
C-312276	GLOC-2_7-9	1.47	0.42	0.61	0.05	0.21	7	491
C-312275	GLOC-2_9-11	1.50	0.44	0.60	0.06	0.22	7	512
C-312258	GLOC-2_11-13	1.48	0.39	0.68	0.05	0.22	6	484
C-312242	GLOC-2_13-15	1.47	0.35	0.72	0.06	0.22	5	467
C-312302	GLOC-2_15-17	1.52	0.41	0.68	0.05	0.25	6	501
C-312245	GLOC-2_17-19	1.56	0.43	0.67	0.04	0.25	6	504
C-312254	GLOC-2_19-21	1.52	0.46	0.59	0.04	0.24	8	539
C-312300	GLOC-2_21-23	1.57	0.44	0.64	0.03	0.25	8	530
C-312247	GLOC-2_23-25	1.51	0.43	0.62	0.03	0.23	6	479
C-312297	GLOC-2_25-27	1.38	0.4	0.58	0.03	0.24	6	475
C-312283	GLOC-2_27-29	1.53	0.44	0.64	0.03	0.22	7	523
C-312266	GLOC-2_29-31	1.44	0.37	0.67	0.03	0.21	6	499
C-312309	GLOC-2_31-33	1.42	0.39	0.62	0.04	0.23	7	496
C-312311	GLOC-2_33-35	1.51	0.44	0.61	0.04	0.27	6	527
C-312307	GLOC-2_35-37	1.50	0.46	0.57	0.04	0.27	7	520
C-312292	GLOC-2_37-39	1.34	0.39	0.56	0.09	0.21	8	443
C-312298	GLOC-2_39-41	1.54	0.44	0.65	0.05	0.24	6	507
C-312253	GLOC-2_41-43	1.49	0.42	0.64	0.06	0.22	7	502
C-312271	GLOC-2_43-45	1.53	0.5	0.56	0.06	0.25	7	533
C-312282	GLOC-2_45-47	1.49	0.46	0.57	0.08	0.26	10	502
C-312310	GLOC-2_47-49	1.48	0.45	0.59	0.05	0.27	8	506
C-312241	GLOC-2_49-51	1.60	0.42	0.72	0.04	0.23	4	507
C-312308	GLOC-2_51-53	1.54	0.39	0.73	0.04	0.24	5	506
C-312286	GLOC-2_53-55	1.52	0.47	0.59	0.05	0.23	7	515
C-312248	GLOC-2_55-57	1.58	0.51	0.57	0.05	0.27	8	540
C-312240	GLOC-2_57-59	1.57	0.56	0.52	0.04	0.26	6	546
C-312280	GLOC-2_59-61	1.37	0.36	0.65	0.03	0.2	5	451
C-312287	GLOC-2_61-63	1.36	0.39	0.59	0.03	0.2	5	454
C-312261	GLOC-2_63-65	1.47	0.44	0.6	0.04	0.25	9	507
C-312289	GLOC-2_65-67	1.37	0.4	0.58	0.04	0.2	8	469
C-312285	GLOC-2_67-69	1.43	0.44	0.57	0.04	0.22	7	489
C-312237	GLOC-2_69-71	1.57	0.51	0.55	0.05	0.24	9	523
C-312304	GLOC-2_71-73	1.54	0.51	0.56	0.04	0.26	8	522
C-312255	GLOC-2_73-75	1.52	0.55	0.49	0.05	0.26	9	524
C-312281	GLOC-2_75-77	1.50	0.52	0.49	0.06	0.24	12	503
C-312306	GLOC-2_77-79	1.61	0.61	0.48	0.06	0.28	11	552
C-312264	GLOC-2_79-81	1.47	0.49	0.54	0.05	0.22	10	492
C-312249	GLOC-2_81-83	1.48	0.38	0.68	0.03	0.22	6	453
C-312265	GLOC-2_83-85	1.47	0.47	0.59	0.04	0.22	6	511
C-312279	GLOC-2_85-87	1.51	0.54	0.5	0.07	0.24	11	534
C-312301	GLOC-2_87-89	1.54	0.5	0.61	0.05	0.25	10	512
C-312259	GLOC-2_89-91	1.56	0.54	0.6	0.05	0.23	9	553

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued
 [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	K %	Mg %	Na %	S %	Ti %	As ppm	Ba ppm
C-312290	GLOC-2_91-93	1.68	0.76	0.41	0.05	0.26	9	532
C-312238	GLOC-2_93-95	1.53	0.53	0.5	0.06	0.27	15	571
C-312303	GLOC-2_95-97	1.44	0.47	0.54	0.05	0.26	7	503
C-312251	GLOC-2_97-99	1.53	0.52	0.52	0.05	0.26	8	506
C-312236	GLOC-2_99-101	1.70	0.62	0.51	0.05	0.25	9	549
C-312293	GLOC-2_101-103	1.61	0.59	0.51	0.04	0.25	9	549
C-312277	GLOC-2_103-105	1.51	0.54	0.48	0.05	0.24	12	516
C-312262	GLOC-2_105-107	1.54	0.52	0.57	0.05	0.25	7	515
C-312246	GLOC-2_107-109	1.53	0.53	0.51	0.06	0.26	7	498
C-312291	GLOC-2_109-111	1.37	0.45	0.48	0.06	0.21	8	481
C-312256	GLOC-2_111-113	1.26	0.36	0.55	0.05	0.2	7	417
C-312299	GLOC-2_113-115	1.40	0.44	0.59	0.04	0.23	7	472
C-312295	GLOC-2_115-117	1.54	0.56	0.5	0.06	0.27	12	542
C-312274	GLOC-2_117-119	1.51	0.56	0.49	0.05	0.24	8	526
C-312250	GLOC-2_119-121	1.58	0.54	0.49	0.05	0.28	10	523
C-312272	GLOC-2_121-123	1.54	0.58	0.41	0.06	0.29	10	556

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Bi ppm	Cd ppm	Co ppm	Cr ppm	Cu ppm	Ga ppm	In ppm
C-312239	GLOC-2_1-3	0.21	0.4	10.6	50	14.8	12.9	0.04
C-312296	GLOC-2_3-5	0.25	0.4	10.3	52	14.9	13.6	0.04
C-312263	GLOC-2_5-7	0.23	0.5	10.9	51	15.2	13.4	0.04
C-312276	GLOC-2_7-9	0.22	0.8	9.9	53	15.7	12.8	0.04
C-312275	GLOC-2_9-11	0.24	0.6	11	53	16.5	13.6	0.04
C-312258	GLOC-2_11-13	0.21	0.6	9.4	50	14.6	12.1	0.04
C-312242	GLOC-2_13-15	0.17	0.5	8.7	40	12.9	10.5	0.04
C-312302	GLOC-2_15-17	0.22	0.6	9.5	47	14.3	12.6	0.04
C-312245	GLOC-2_17-19	0.21	0.6	10.2	50	15.6	12.3	0.04
C-312254	GLOC-2_19-21	0.26	1.2	9.6	61	18.4	14.1	0.05
C-312300	GLOC-2_21-23	0.26	0.7	11	49	16.9	13.8	0.05
C-312247	GLOC-2_23-25	0.93	0.8	10.1	47	15.4	12.8	0.04
C-312297	GLOC-2_25-27	0.25	0.8	10.2	51	14.8	12.8	0.04
C-312283	GLOC-2_27-29	0.27	0.7	9.9	52	16.3	13.5	0.04
C-312266	GLOC-2_29-31	0.23	0.7	8.5	49	14.5	11.8	0.04
C-312309	GLOC-2_31-33	0.26	1.3	8.5	49	16.5	12.5	0.04
C-312311	GLOC-2_33-35	0.27	1.7	9.4	60	17.7	14	0.05
C-312307	GLOC-2_35-37	0.29	1.6	10.2	63	17.3	14.1	0.05
C-312292	GLOC-2_37-39	0.26	1.2	10.6	49	16.3	12.8	0.04
C-312298	GLOC-2_39-41	0.25	1.1	8.6	51	16	12.6	0.04
C-312253	GLOC-2_41-43	0.23	0.8	10.2	52	17	13	0.04
C-312271	GLOC-2_43-45	0.3	0.9	11.6	59	20.6	14.9	0.05
C-312282	GLOC-2_45-47	0.32	2	11.7	58	19	14.8	0.05
C-312310	GLOC-2_47-49	0.29	1	11	56	16.9	14.3	0.05
C-312241	GLOC-2_49-51	0.2	0.4	9.4	43	14.2	11.5	0.04
C-312308	GLOC-2_51-53	0.23	0.4	9.4	47	13.1	12.2	0.04
C-312286	GLOC-2_53-55	0.25	0.6	10.2	57	17.1	13.8	0.05
C-312248	GLOC-2_55-57	0.25	1.5	11	55	18.1	13.9	0.05
C-312240	GLOC-2_57-59	0.27	2.2	10.5	58	19.7	14.6	0.05
C-312280	GLOC-2_59-61	0.2	0.7	9.4	43	12.7	11.4	0.04
C-312287	GLOC-2_61-63	0.21	1	9	46	13.7	11.6	0.06
C-312261	GLOC-2_63-65	0.26	1.3	10.2	53	15.5	13.3	0.05
C-312289	GLOC-2_65-67	0.3	1.3	9.3	64	17.9	12.5	0.05
C-312285	GLOC-2_67-69	0.24	1.4	9	56	16.3	12.7	0.06
C-312237	GLOC-2_69-71	0.26	1.5	11.3	59	23	14.6	0.07
C-312304	GLOC-2_71-73	0.27	1	11.2	57	18.7	14.9	0.06
C-312255	GLOC-2_73-75	0.29	2.1	12.2	68	21	15.7	0.08
C-312281	GLOC-2_75-77	0.29	3.1	11.9	65	21	15.2	0.09
C-312306	GLOC-2_77-79	0.31	1.3	12.9	69	21.2	17.1	0.07
C-312264	GLOC-2_79-81	0.25	1.2	9.7	56	17.6	14	0.06
C-312249	GLOC-2_81-83	0.2	0.8	8	37	13.6	11.1	0.05
C-312265	GLOC-2_83-85	0.24	1.4	8.5	55	17.4	13.3	0.06
C-312279	GLOC-2_85-87	0.31	4.1	10.5	61	23.3	15.2	0.17
C-312301	GLOC-2_87-89	0.27	0.8	12.4	54	17.2	14.4	0.07
C-312259	GLOC-2_89-91	0.25	0.6	10.1	55	17.8	13.7	0.05

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued
 [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Bi ppm	Cd ppm	Co ppm	Cr ppm	Cu ppm	Ga ppm	In ppm
C-312290	GLOC-2_91-93	0.35	0.8	13	65	26.7	18.2	0.07
C-312238	GLOC-2_93-95	0.28	4	10.2	59	21.6	14.8	0.1
C-312303	GLOC-2_95-97	0.31	2.9	9.5	58	19.7	14.5	0.08
C-312251	GLOC-2_97-99	0.31	2.6	11.5	62	20.4	15	0.09
C-312236	GLOC-2_99-101	0.3	1.6	14.6	63	26.5	15.9	0.11
C-312293	GLOC-2_101-103	0.31	1.2	11.6	65	21	17	0.12
C-312277	GLOC-2_103-105	0.33	2.7	11.5	69	24.2	16	0.22
C-312262	GLOC-2_105-107	0.3	1.1	11.3	63	18	14.7	0.07
C-312246	GLOC-2_107-109	0.25	1	10.8	60	19.1	14.3	0.08
C-312291	GLOC-2_109-111	0.24	1	10	58	16.2	13.4	0.06
C-312256	GLOC-2_111-113	0.21	1	9.7	47	14.2	11.2	0.09
C-312299	GLOC-2_113-115	0.26	1.2	9.1	52	16.8	12.3	0.13
C-312295	GLOC-2_115-117	0.34	2.4	12.4	63	21.5	15.9	0.13
C-312274	GLOC-2_117-119	0.3	1.4	13	66	22.5	16.1	0.08
C-312250	GLOC-2_119-121	0.31	3	12.1	66	25.7	16.7	0.1
C-312272	GLOC-2_121-123	0.39	3.7	11.5	74	29.6	17.1	0.19

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Li ppm	Mn ppm	Ni ppm	P ppm	Pb ppm	Rb ppm	Sr ppm
C-312239	GLOC-2_1-3	31	521	22.7	630	19.6	94.1	113
C-312296	GLOC-2_3-5	30	472	21.8	580	20.2	89	113
C-312263	GLOC-2_5-7	30	470	21.3	610	20.8	87.2	113
C-312276	GLOC-2_7-9	29	456	22.3	660	21.7	84.5	110
C-312275	GLOC-2_9-11	31	475	22.4	660	22.9	89.8	112
C-312258	GLOC-2_11-13	27	404	20.1	600	19.6	80.8	108
C-312242	GLOC-2_13-15	25	320	17.6	470	18.7	78.6	101
C-312302	GLOC-2_15-17	30	398	20.6	570	20.4	84.3	110
C-312245	GLOC-2_17-19	31	434	22.5	560	22.2	87	108
C-312254	GLOC-2_19-21	33	522	25.6	710	27.8	95.8	112
C-312300	GLOC-2_21-23	32	446	22.7	580	24.2	94.6	109
C-312247	GLOC-2_23-25	31	467	21.5	640	27.0	91.5	102
C-312297	GLOC-2_25-27	29	422	22.2	600	24.6	82.3	100
C-312283	GLOC-2_27-29	31	478	21.7	620	24.5	87.4	110
C-312266	GLOC-2_29-31	27	414	19.5	510	24.9	79.4	107
C-312309	GLOC-2_31-33	30	408	20.3	620	26.7	83.3	105
C-312311	GLOC-2_33-35	33	450	24.1	680	28.8	91.4	111
C-312307	GLOC-2_35-37	33	413	24.4	700	28.7	92.1	106
C-312292	GLOC-2_37-39	29	402	22.3	690	26.4	82.8	101
C-312298	GLOC-2_39-41	32	689	22.5	670	23.2	88.3	110
C-312253	GLOC-2_41-43	30	506	22.2	570	22.8	85.2	109
C-312271	GLOC-2_43-45	34	804	25.8	730	25.0	96.8	116
C-312282	GLOC-2_45-47	34	526	26.6	670	30.2	92.9	109
C-312310	GLOC-2_47-49	34	470	24.3	580	26.4	91.4	113
C-312241	GLOC-2_49-51	29	740	19.8	470	19.5	82.1	114
C-312308	GLOC-2_51-53	28	603	19.7	480	19.1	81.1	115
C-312286	GLOC-2_53-55	33	721	24.2	590	23.7	92	119
C-312248	GLOC-2_55-57	35	355	25.6	560	27.6	94.8	114
C-312240	GLOC-2_57-59	38	335	28	630	29.1	101	114
C-312280	GLOC-2_59-61	24	495	17.8	490	21.0	76.7	104
C-312287	GLOC-2_61-63	26	480	19.7	500	23.1	75.8	102
C-312261	GLOC-2_63-65	30	598	22.1	630	25.5	85.2	113
C-312289	GLOC-2_65-67	28	404	24.2	520	24.6	81.9	103
C-312285	GLOC-2_67-69	31	488	23.7	560	26.5	85	109
C-312237	GLOC-2_69-71	36	635	27.6	650	27.9	98.2	115
C-312304	GLOC-2_71-73	36	649	26.7	660	24.7	96	116
C-312255	GLOC-2_73-75	37	787	30	680	29.7	99.9	120
C-312281	GLOC-2_75-77	36	649	29.2	650	34.5	97.7	117
C-312306	GLOC-2_77-79	40	715	31.2	690	25.7	109	126
C-312264	GLOC-2_79-81	32	569	24.7	570	24.5	87.1	116
C-312249	GLOC-2_81-83	26	577	17.8	440	20.9	81	109
C-312265	GLOC-2_83-85	30	461	22.4	550	21.6	84.5	119
C-312279	GLOC-2_85-87	36	597	27.2	740	39.1	94.6	126
C-312301	GLOC-2_87-89	34	581	24.9	550	22.7	90.6	132
C-312259	GLOC-2_89-91	31	560	22.2	590	21.2	90.8	129

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued
 [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Li ppm	Mn ppm	Ni ppm	P ppm	Pb ppm	Rb ppm	Sr ppm
C-312290	GLOC-2_91-93	43	932	28.4	690	28.3	114	130
C-312238	GLOC-2_93-95	38	522	27.1	750	40.6	98.5	122
C-312303	GLOC-2_95-97	35	541	24.8	700	35.4	90.9	115
C-312251	GLOC-2_97-99	36	606	29.4	710	33.6	101	115
C-312236	GLOC-2_99-101	41	981	32.5	760	27.9	110	123
C-312293	GLOC-2_101-103	39	1,160	29.1	700	26.1	106	128
C-312277	GLOC-2_103-105	37	670	29.8	610	33.2	101	118
C-312262	GLOC-2_105-107	34	539	27	560	24.7	92.7	114
C-312246	GLOC-2_107-109	36	537	28.1	560	27.0	98.4	111
C-312291	GLOC-2_109-111	31	518	26.5	530	24.3	85.4	104
C-312256	GLOC-2_111-113	27	466	20.2	470	23.0	70.3	98.1
C-312299	GLOC-2_113-115	30	576	23.4	520	23.3	81.4	116
C-312295	GLOC-2_115-117	37	848	30	830	29.3	102	124
C-312274	GLOC-2_117-119	38	508	29.9	690	29.8	102	111
C-312250	GLOC-2_119-121	42	485	29.4	750	45.0	110	111
C-312272	GLOC-2_121-123	44	759	31.8	980	41.2	113	112

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued
 [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	V ppm	Zn ppm
C-312239	GLOC-2_1-3	77	95
C-312296	GLOC-2_3-5	78	113
C-312263	GLOC-2_5-7	78	157
C-312276	GLOC-2_7-9	80	197
C-312275	GLOC-2_9-11	84	152
C-312258	GLOC-2_11-13	72	167
C-312242	GLOC-2_13-15	62	125
C-312302	GLOC-2_15-17	76	161
C-312245	GLOC-2_17-19	77	131
C-312254	GLOC-2_19-21	89	189
C-312300	GLOC-2_21-23	80	132
C-312247	GLOC-2_23-25	75	153
C-312297	GLOC-2_25-27	78	180
C-312283	GLOC-2_27-29	79	142
C-312266	GLOC-2_29-31	72	113
C-312309	GLOC-2_31-33	75	231
C-312311	GLOC-2_33-35	88	289
C-312307	GLOC-2_35-37	87	279
C-312292	GLOC-2_37-39	76	283
C-312298	GLOC-2_39-41	75	234
C-312253	GLOC-2_41-43	78	185
C-312271	GLOC-2_43-45	94	196
C-312282	GLOC-2_45-47	87	475
C-312310	GLOC-2_47-49	88	271
C-312241	GLOC-2_49-51	70	113
C-312308	GLOC-2_51-53	71	99
C-312286	GLOC-2_53-55	88	141
C-312248	GLOC-2_55-57	86	372
C-312240	GLOC-2_57-59	93	453
C-312280	GLOC-2_59-61	67	162
C-312287	GLOC-2_61-63	69	243
C-312261	GLOC-2_63-65	81	315
C-312289	GLOC-2_65-67	76	281
C-312285	GLOC-2_67-69	81	327
C-312237	GLOC-2_69-71	90	327
C-312304	GLOC-2_71-73	87	236
C-312255	GLOC-2_73-75	98	439
C-312281	GLOC-2_75-77	95	563
C-312306	GLOC-2_77-79	99	339
C-312264	GLOC-2_79-81	86	288
C-312249	GLOC-2_81-83	64	166
C-312265	GLOC-2_83-85	85	393
C-312279	GLOC-2_85-87	93	1,000
C-312301	GLOC-2_87-89	85	194
C-312259	GLOC-2_89-91	82	131

Table 1–4. Core sediment sample analyses for GLOC-2, Neosho River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued
 [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	V ppm	Zn ppm
C-312290	GLOC-2_91-93	112	170
C-312238	GLOC-2_93-95	94	649
C-312303	GLOC-2_95-97	88	536
C-312251	GLOC-2_97-99	91	464
C-312236	GLOC-2_99-101	100	364
C-312293	GLOC-2_101-103	102	212
C-312277	GLOC-2_103-105	101	420
C-312262	GLOC-2_105-107	88	219
C-312246	GLOC-2_107-109	93	218
C-312291	GLOC-2_109-111	85	227
C-312256	GLOC-2_111-113	71	215
C-312299	GLOC-2_113-115	75	265
C-312295	GLOC-2_115-117	96	563
C-312274	GLOC-2_117-119	101	244
C-312250	GLOC-2_119-121	101	438
C-312272	GLOC-2_121-123	112	634

Table 1–5. Core sediment sample analyses for GLOC-3, Spring River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon. ([Click to see Excel file.](#))
 [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million; n/a, not applicable]

Lab no.	Field no.	Depth in inches	Depth in cm	Total C %	Carbonate C %	Organic C %	Al %	Ca %	Fe %
C-312208	GLOC-3_0-2	2	5	0.8	0.06	0.74	1.78	0.31	1.31
C-312225	GLOC-3_2-4	4	10	2.09	0.11	1.98	4.12	0.68	2.04
C-312231	GLOC-3_4-6	6	15	1.27	0.08	1.19	2.71	0.45	1.61
C-312228	GLOC-3_6-8	8	20	1.07	0.06	1.01	2.40	0.36	1.49
C-312224	GLOC-3_8-10	10	25	1.28	0.04	1.24	3.11	0.38	1.69
C-312198	GLOC-3_10-12	12	30	1.55	0.02	1.53	3.48	0.37	1.91
C-312200	GLOC-3_12-14	14	35	1.75	0.03	1.72	4.70	0.48	2.36
C-312192	GLOC-3_14-16	16	40	1.49	0.08	1.41	3.62	0.53	1.82
C-312188	GLOC-3_16-18	18	45	1.35	0.07	1.28	3.25	0.43	1.70
C-312191	GLOC-3_18-20	20	50	1.48	0.04	1.44	3.83	0.40	1.95
C-312218	GLOC-3_20-22	22	55	n/a	0.05	n/a	4.41	0.42	2.07
C-312210	GLOC-3_22-24	24	60	1.52	0.01	1.51	4.71	0.36	2.13
C-312211	GLOC-3_24-26	26	65	1.44	0.01	1.43	4.42	0.33	2.09
C-312204	GLOC-3_26-28	28	70	1.47	0.01	1.46	4.51	0.33	2.08
C-312216	GLOC-3_28-30	30	75	1.56	0.01	1.55	4.74	0.34	2.15
C-312197	GLOC-3_30-32	32	80	1.58	<0.003	1.58	4.83	0.35	2.23
C-312215	GLOC-3_32-34	34	85	1.53	0.01	1.52	4.85	0.34	2.23
C-312212	GLOC-3_34-36	36	90	1.48	0.01	1.47	4.90	0.34	2.28
C-312189	GLOC-3_36-38	38	95	1.46	0.02	1.44	4.88	0.34	2.37
C-312226	GLOC-3_38-40	40	100	1.47	0.02	1.45	5.08	0.35	2.50
C-312194	GLOC-3_40-42	42	105	1.55	0.02	1.53	4.82	0.33	2.32
C-312230	GLOC-3_42-44	44	110	1.45	0.04	1.41	5.27	0.36	2.49
C-312219	GLOC-3_44-46	46	115	1.42	0.01	1.41	4.60	0.33	2.27
C-312223	GLOC-3_46-48	48	120	1.39	0.01	1.38	4.68	0.33	2.21
C-312203	GLOC-3_48-50	50	125	1.43	0.03	1.40	4.52	0.32	2.06
C-312195	GLOC-3_50-52	52	130	1.31	0.01	1.30	4.16	0.28	1.72
C-312209	GLOC-3_52-54	54	135	1.1	0.07	1.03	4.27	0.31	1.94
C-312205	GLOC-3_54-56	56	140	0.92	0.02	0.90	4.25	0.27	1.66
C-312206	GLOC-3_56-58	58	145	0.85	<0.003	0.85	4.02	0.26	1.44
C-312214	GLOC-3_58-60	60	150	0.65	0.01	0.64	3.69	0.22	1.35
C-312213	GLOC-3_60-62	62	155	0.66	0.01	0.65	3.82	0.23	1.38
C-312196	GLOC-3_62-64	64	160	0.66	0.01	0.65	3.83	0.23	1.35
C-312222	GLOC-3_64-66	66	165	0.59	0.01	0.58	3.86	0.23	1.39
C-312221	GLOC-3_66-68	68	170	0.54	0.02	0.52	3.93	0.22	1.43
C-312227	GLOC-3_68-70	70	175	0.49	0.02	0.47	3.79	0.21	1.30
C-312202	GLOC-3_70-72	72	180	0.49	0.01	0.48	3.85	0.21	1.25
C-312220	GLOC-3_72-74	74	185	0.43	<0.003	0.43	4.02	0.24	1.34

Table 1–5. Core sediment sample analyses for GLOC-3, Spring River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million; n/a, not applicable]

Lab no.	Field no.	K %	Mg %	Na %	S %	Ti %	As ppm	Ba ppm	Bi ppm
C-312208	GLOC-3_0-2	0.43	0.11	0.14	0.05	0.09	3	169	0.11
C-312225	GLOC-3_2-4	1.02	0.24	0.32	0.1	0.21	6	386	0.23
C-312231	GLOC-3_4-6	0.68	0.16	0.22	0.07	0.15	4	253	0.15
C-312228	GLOC-3_6-8	0.63	0.15	0.23	0.05	0.13	3	236	0.14
C-312224	GLOC-3_8-10	0.84	0.18	0.30	0.05	0.2	5	312	0.17
C-312198	GLOC-3_10-12	0.85	0.19	0.28	0.05	0.19	5	324	0.19
C-312200	GLOC-3_12-14	1.11	0.27	0.34	0.09	0.24	7	432	0.27
C-312192	GLOC-3_14-16	0.94	0.21	0.33	0.11	0.21	6	343	0.20
C-312188	GLOC-3_16-18	0.84	0.18	0.29	0.12	0.18	5	315	0.19
C-312191	GLOC-3_18-20	0.95	0.22	0.31	0.1	0.21	5	357	0.24
C-312218	GLOC-3_20-22	1.09	0.25	0.35	0.12	0.23	9	406	0.28
C-312210	GLOC-3_22-24	1.15	0.26	0.36	0.09	0.24	8	429	0.36
C-312211	GLOC-3_24-26	1.08	0.24	0.34	0.06	0.24	5	413	0.37
C-312204	GLOC-3_26-28	1.11	0.25	0.35	0.06	0.24	5	438	0.33
C-312216	GLOC-3_28-30	1.15	0.27	0.35	0.07	0.24	5	446	0.33
C-312197	GLOC-3_30-32	1.16	0.27	0.37	0.05	0.25	6	468	0.38
C-312215	GLOC-3_32-34	1.15	0.27	0.35	0.05	0.25	6	452	0.37
C-312212	GLOC-3_34-36	1.15	0.28	0.34	0.05	0.25	7	440	0.38
C-312189	GLOC-3_36-38	1.14	0.28	0.33	0.04	0.26	7	438	0.34
C-312226	GLOC-3_38-40	1.2	0.29	0.34	0.04	0.3	7	450	0.35
C-312194	GLOC-3_40-42	1.16	0.27	0.33	0.05	0.25	7	443	0.34
C-312230	GLOC-3_42-44	1.30	0.30	0.38	0.05	0.25	7	449	0.31
C-312219	GLOC-3_44-46	1.17	0.26	0.35	0.04	0.25	7	441	0.27
C-312223	GLOC-3_46-48	1.22	0.26	0.36	0.04	0.25	7	460	0.27
C-312203	GLOC-3_48-50	1.20	0.24	0.37	0.03	0.25	7	468	0.26
C-312195	GLOC-3_50-52	1.18	0.22	0.39	0.03	0.26	5	490	0.21
C-312209	GLOC-3_52-54	1.21	0.22	0.41	0.03	0.28	5	514	0.20
C-312205	GLOC-3_54-56	1.14	0.22	0.38	0.02	0.24	4	536	0.20
C-312206	GLOC-3_56-58	1.11	0.21	0.37	0.02	0.22	3	497	0.18
C-312214	GLOC-3_58-60	1.06	0.19	0.35	0.02	0.23	3	437	0.17
C-312213	GLOC-3_60-62	1.09	0.20	0.36	0.02	0.23	3	439	0.20
C-312196	GLOC-3_62-64	1.10	0.19	0.36	0.02	0.25	3	452	0.17
C-312222	GLOC-3_64-66	1.12	0.20	0.36	0.02	0.22	3	444	0.19
C-312221	GLOC-3_66-68	1.13	0.20	0.37	0.02	0.25	3	438	0.19
C-312227	GLOC-3_68-70	1.09	0.19	0.35	0.01	0.23	3	419	0.17
C-312202	GLOC-3_70-72	1.10	0.20	0.34	0.01	0.23	3	418	0.18
C-312220	GLOC-3_72-74	1.14	0.21	0.35	0.01	0.24	3	431	0.18

Table 1–5. Core sediment sample analyses for GLOC-3, Spring River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million; n/a, not applicable]

Lab no.	Field no.	Cd ppm	Co ppm	Cr ppm	Cu ppm	Ga ppm	In ppm	Li ppm	Mn ppm
C-312208	GLOC-3_0-2	6.1	6.8	25	10.7	4.23	<0.02	13	357
C-312225	GLOC-3_2-4	12.9	12.5	44	23.7	9.74	0.04	27	796
C-312231	GLOC-3_4-6	8.6	8.7	33	15.2	6.83	0.04	19	477
C-312228	GLOC-3_6-8	7.1	8.3	29	14	5.6	0.02	17	344
C-312224	GLOC-3_8-10	8.2	9.9	34	16.6	6.92	0.03	21	404
C-312198	GLOC-3_10-12	8.1	10.4	38	18.8	7.72	0.03	23	477
C-312200	GLOC-3_12-14	12.2	14.8	48	22.9	11.1	0.04	30	573
C-312192	GLOC-3_14-16	120	11.4	39	17.6	8.32	0.03	24	509
C-312188	GLOC-3_16-18	11.0	9.9	36	18.5	7.52	0.03	21	481
C-312191	GLOC-3_18-20	14.0	12.3	42	22.6	9.01	0.04	25	528
C-312218	GLOC-3_20-22	15.4	17.7	46	24.9	10.4	0.04	28	573
C-312210	GLOC-3_22-24	17.8	14.9	50	36.1	11.2	0.06	30	539
C-312211	GLOC-3_24-26	21.1	13.6	51	38.3	10.3	0.05	28	513
C-312204	GLOC-3_26-28	23.3	13.4	52	31.2	10.6	0.05	28	504
C-312216	GLOC-3_28-30	26.6	13.7	55	32.3	10.8	0.05	29	530
C-312197	GLOC-3_30-32	31.4	14.4	58	40.5	11.4	0.07	31	603
C-312215	GLOC-3_32-34	28.7	13.9	59	39	11.6	0.06	30	522
C-312212	GLOC-3_34-36	29.4	13.9	59	38.7	11.6	0.06	31	472
C-312189	GLOC-3_36-38	20.5	14.6	56	38.8	11.9	0.06	31	576
C-312226	GLOC-3_38-40	23.7	14.7	56	43.7	11.9	0.06	33	627
C-312194	GLOC-3_40-42	24.4	14.3	51	40	11.4	0.05	30	513
C-312230	GLOC-3_42-44	23.5	14.5	53	38.2	11.3	0.05	33	634
C-312219	GLOC-3_44-46	19.7	13.7	46	27.6	11	0.04	28	822
C-312223	GLOC-3_46-48	20.0	12.9	46	25.1	10.9	0.04	28	778
C-312203	GLOC-3_48-50	14.5	11.8	43	24.2	10.4	0.04	28	720
C-312195	GLOC-3_50-52	8.0	9.8	40	17.6	9.65	0.03	27	579
C-312209	GLOC-3_52-54	4.8	10.3	43	17.9	9.43	0.03	27	1,210
C-312205	GLOC-3_54-56	2.0	8.7	42	17.5	9.91	0.04	26	662
C-312206	GLOC-3_56-58	0.6	7.8	39	15.2	8.92	0.03	24	443
C-312214	GLOC-3_58-60	0.4	6.4	37	13.4	8.11	0.02	23	554
C-312213	GLOC-3_60-62	0.5	7.0	36	16.5	8.59	0.03	23	551
C-312196	GLOC-3_62-64	0.3	6.7	35	13.6	8.34	0.03	23	516
C-312222	GLOC-3_64-66	0.3	7.8	37	14.7	8.94	0.03	23	642
C-312221	GLOC-3_66-68	0.3	8.7	37	15.9	8.52	0.03	24	663
C-312227	GLOC-3_68-70	0.2	9.2	38	14.5	8.64	0.03	23	574
C-312202	GLOC-3_70-72	0.3	9.8	36	14.5	8.82	0.03	24	609
C-312220	GLOC-3_72-74	0.3	9.0	38	17.4	9.1	0.03	24	714

Table 1–5. Core sediment sample analyses for GLOC-3, Spring River upstream from Grand Lake O' the Cherokees. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million; n/a, not applicable]

Lab no.	Field no.	Ni ppm	P ppm	Pb ppm	Rb ppm	Sr ppm	V ppm	Zn ppm
C-312208	GLOC-3_0-2	12.5	530	60.5	28.3	28.9	31	1,020
C-312225	GLOC-3_2-4	21.1	1,000	130	68.5	56.9	62	1,820
C-312231	GLOC-3_4-6	15.7	670	69.3	43.6	40.2	43	1,280
C-312228	GLOC-3_6-8	14.6	610	76.5	39.2	36.1	38	1,250
C-312224	GLOC-3_8-10	17.2	640	89.9	50.6	46.4	46	1,440
C-312198	GLOC-3_10-12	19.7	810	95.1	55.1	49.4	52	1,380
C-312200	GLOC-3_12-14	25.1	1,050	109	79.3	60.5	71	1,610
C-312192	GLOC-3_14-16	21.4	710	103	56.9	51.8	51	1,630
C-312188	GLOC-3_16-18	20.4	650	98.4	52.2	45.7	48	1,550
C-312191	GLOC-3_18-20	24.2	770	122	60.9	51.1	58	1,710
C-312218	GLOC-3_20-22	29.7	880	143	73.8	56.5	66	1,910
C-312210	GLOC-3_22-24	28.1	900	196	78.1	59.8	70	2,180
C-312211	GLOC-3_24-26	27	920	226	71.1	57.7	65	2,380
C-312204	GLOC-3_26-28	28.5	960	176	74.2	58.5	67	2,470
C-312216	GLOC-3_28-30	30.8	1,130	176	75.4	59.9	71	2,640
C-312197	GLOC-3_30-32	32.9	1,430	173	79.8	62.8	71	2,980
C-312215	GLOC-3_32-34	30.8	1,200	177	78.9	60.9	74	3,120
C-312212	GLOC-3_34-36	30	950	172	79.3	60.5	75	3,240
C-312189	GLOC-3_36-38	30.3	770	180	81.9	60.2	77	3,160
C-312226	GLOC-3_38-40	31.3	740	207	84.1	63.3	76	3,300
C-312194	GLOC-3_40-42	30.6	640	248	78.5	59.9	73	3,340
C-312230	GLOC-3_42-44	31.4	640	208	80.5	67.2	77	3,250
C-312219	GLOC-3_44-46	27.8	600	169	80	59.1	72	2,810
C-312223	GLOC-3_46-48	28.1	590	157	82.7	60.5	73	2,400
C-312203	GLOC-3_48-50	25.5	580	128	84.3	62.2	70	1,800
C-312195	GLOC-3_50-52	21.7	480	84.6	78.7	63.4	66	1,060
C-312209	GLOC-3_52-54	19.9	460	48.6	79.2	68.6	63	606
C-312205	GLOC-3_54-56	17.4	390	25.2	81.7	65.9	65	297
C-312206	GLOC-3_56-58	15.1	370	20	75.8	60.5	60	154
C-312214	GLOC-3_58-60	13.8	310	18.3	66.9	55.9	52	114
C-312213	GLOC-3_60-62	13.8	310	19.7	71.6	57.3	55	113
C-312196	GLOC-3_62-64	13.3	290	19.1	68	57.8	54	100
C-312222	GLOC-3_64-66	14	280	19.9	71.3	56.2	54	102
C-312221	GLOC-3_66-68	14.6	260	20.2	71.1	58.8	54	100
C-312227	GLOC-3_68-70	14.4	240	18.7	73.9	55	52	97
C-312202	GLOC-3_70-72	14.3	230	17.4	74.7	54.9	52	100
C-312220	GLOC-3_72-74	15.1	220	18.8	75.1	56.3	54	103

Table 1–6. Core sediment sample analyses for GLOC-7, Grand Lake O' the Cherokees 4 kilometers downstream from Highway 2 bridge. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.

([Click to see Excel file.](#))

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Depth in inches	Depth in cm	Total C %	Carbonate C %	Organic C %	Al %	Ca %	Fe %
C-312182	GLOC-7_0-2	2	5	0.90	0.06	0.84	4.63	0.59	1.89
C-312172	GLOC-7_2-4	4	10	0.92	0.04	0.88	4.56	0.51	1.87
C-312159	GLOC-7_4-6	6	15	0.73	0.01	0.72	4.96	0.40	1.95
C-312160	GLOC-7_6-8	8	20	1.04	0.01	1.03	5.76	0.46	2.54
C-312169	GLOC-7_8-10	10	25	0.96	0.01	0.95	5.79	0.47	2.57
C-312154	GLOC-7_10-12	12	30	0.71	0.01	0.70	5.21	0.43	2.14
C-312176	GLOC-7_12-14	14	35	0.80	0.01	0.79	5.07	0.43	2.09
C-312183	GLOC-7_14-16	16	40	0.88	0.02	0.86	5.44	0.48	2.40
C-312162	GLOC-7_17-18	18	45	0.72	0.01	0.71	4.97	0.42	1.94
C-312151	GLOC-7_18-20	20	50	0.91	0.05	0.86	5.75	0.62	2.44
C-312155	GLOC-7_20-22	22	55	0.93	0.03	0.90	5.31	0.50	2.18
C-312158	GLOC-7_22-24	24	60	0.79	<0.003	0.79	5.01	0.47	2.03
C-312179	GLOC-7_24-26	26	65	1.09	0.05	1.04	5.76	0.59	2.49
C-312167	GLOC-7_26-28	28	70	1.28	0.09	1.19	5.66	0.64	2.53
C-312178	GLOC-7_28-30	30	75	1.00	0.01	0.99	5.09	0.36	2.17
C-312170	GLOC-7_30-32	32	80	0.84	<0.003	0.84	3.91	0.23	1.57
C-312157	GLOC-7_32-34	34	85	0.79	<0.003	0.79	3.84	0.20	1.55
C-312163	GLOC-7_34-36	36	90	0.83	<0.003	0.83	3.96	0.21	1.60
C-312177	GLOC-7_36-38	38	95	0.82	<0.003	0.82	3.98	0.20	1.65
C-312168	GLOC-7_38-40	40	100	0.80	<0.003	0.80	4.02	0.20	1.73
C-312166	GLOC-7_40-42	42	105	0.75	<0.003	0.75	3.77	0.18	1.59
C-312173	GLOC-7_42-44	44	110	0.76	<0.003	0.76	4.11	0.19	1.80
C-312180	GLOC-7_44-46	46	115	0.73	<0.003	0.73	4.19	0.20	1.84
C-312171	GLOC-7_46-48	48	120	0.73	<0.003	0.73	4.29	0.20	1.74
C-312156	GLOC-7_48-50	50	125	0.64	<0.003	0.64	4.30	0.19	1.83
C-312174	GLOC-7_50-52	52	130	0.6	<0.003	0.60	4.39	0.18	1.97
C-312161	GLOC-7_52-54	54	135	0.55	<0.003	0.55	4.49	0.19	2.03

Table 1–6. Core sediment sample analyses for GLOC-7, Grand Lake O' the Cherokees 4 kilometers downstream from Highway 2 bridge. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	K %	Mg %	Na %	S %	Ti %	As ppm	Ba ppm	Bi ppm
C-312182	GLOC-7_0-2	1.42	0.34	0.66	0.06	0.22	6	447	0.18
C-312172	GLOC-7_2-4	1.37	0.34	0.62	0.06	0.2	6	427	0.18
C-312159	GLOC-7_4-6	1.49	0.36	0.68	0.04	0.22	5	474	0.20
C-312160	GLOC-7_6-8	1.56	0.47	0.58	0.04	0.25	8	511	0.25
C-312169	GLOC-7_8-10	1.60	0.48	0.60	0.03	0.25	6	521	0.26
C-312154	GLOC-7_10-12	1.57	0.41	0.69	0.02	0.22	5	501	0.21
C-312176	GLOC-7_12-14	1.51	0.38	0.70	0.02	0.23	5	491	0.21
C-312183	GLOC-7_14-16	1.54	0.44	0.62	0.02	0.23	6	494	0.22
C-312162	GLOC-7_17-18	1.49	0.38	0.68	0.03	0.22	6	486	0.19
C-312151	GLOC-7_18-20	1.67	0.50	0.66	0.05	0.24	7	531	0.24
C-312155	GLOC-7_20-22	1.48	0.40	0.63	0.04	0.2	6	484	0.23
C-312158	GLOC-7_22-24	1.46	0.38	0.66	0.04	0.21	6	465	0.20
C-312179	GLOC-7_24-26	1.57	0.47	0.60	0.05	0.25	7	509	0.25
C-312167	GLOC-7_26-28	1.55	0.48	0.59	0.05	0.25	7	505	0.26
C-312178	GLOC-7_28-30	1.43	0.40	0.51	0.04	0.23	6	475	0.21
C-312170	GLOC-7_30-32	1.14	0.25	0.38	0.05	0.21	4	413	0.17
C-312157	GLOC-7_32-34	1.10	0.24	0.34	0.03	0.19	4	396	0.16
C-312163	GLOC-7_34-36	1.10	0.25	0.35	0.03	0.2	4	410	0.17
C-312177	GLOC-7_36-38	1.11	0.25	0.34	0.02	0.2	5	400	0.17
C-312168	GLOC-7_38-40	1.09	0.25	0.32	0.02	0.2	5	407	0.17
C-312166	GLOC-7_40-42	1.02	0.23	0.29	0.02	0.2	4	390	0.16
C-312173	GLOC-7_42-44	1.10	0.25	0.31	0.02	0.2	5	407	0.17
C-312180	GLOC-7_44-46	1.11	0.26	0.30	0.02	0.22	6	414	0.18
C-312171	GLOC-7_46-48	1.12	0.26	0.30	0.03	0.2	4	432	0.18
C-312156	GLOC-7_48-50	1.10	0.26	0.29	0.02	0.18	4	428	0.18
C-312174	GLOC-7_50-52	1.12	0.27	0.28	0.02	0.22	5	429	0.18
C-312161	GLOC-7_52-54	1.13	0.28	0.28	0.02	0.22	6	432	0.19

Table 1–6. Core sediment sample analyses for GLOC-7, Grand Lake O' the Cherokees 4 kilometers downstream from Highway 2 bridge. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Cd ppm	Co ppm	Cr ppm	Cu ppm	Ga ppm	In ppm	Li ppm	Mn ppm
C-312182	GLOC-7_0-2	1.8	9.1	40	17.7	10.0	0.04	25	384
C-312172	GLOC-7_2-4	2.2	9.5	40	15.8	10.7	0.03	25	366
C-312159	GLOC-7_4-6	1.4	8.2	41	16.3	10.8	0.04	26	385
C-312160	GLOC-7_6-8	4.6	11.2	55	23.8	13.2	0.05	34	612
C-312169	GLOC-7_8-10	2.6	11.1	53	23.0	13.8	0.04	33	598
C-312154	GLOC-7_10-12	1.3	8.3	44	18.0	11.2	0.04	28	489
C-312176	GLOC-7_12-14	2.2	8.7	43	16.5	11.6	0.04	27	458
C-312183	GLOC-7_14-16	1.1	9.6	68	18.9	12.5	0.05	30	730
C-312162	GLOC-7_17-18	0.7	8.4	43	14.2	11.2	0.04	27	403
C-312151	GLOC-7_18-20	1.0	10.0	54	17.0	12.1	0.05	33	636
C-312155	GLOC-7_20-22	2.4	9.0	49	17.5	11.8	0.04	29	565
C-312158	GLOC-7_22-24	1.6	8.7	44	19.0	10.4	0.04	28	582
C-312179	GLOC-7_24-26	3.7	10.8	55	20.3	13.6	0.05	33	678
C-312167	GLOC-7_26-28	4.3	11.4	54	20.9	13.3	0.05	33	1,000
C-312178	GLOC-7_28-30	1.5	10.0	46	18.0	11.8	0.04	29	436
C-312170	GLOC-7_30-32	0.6	8.0	38	15.2	8.89	0.03	23	302
C-312157	GLOC-7_32-34	0.3	7.0	38	13.5	8.39	0.03	22	323
C-312163	GLOC-7_34-36	0.4	7.5	37	14.3	8.94	0.03	23	338
C-312177	GLOC-7_36-38	0.4	7.5	36	14.7	9.15	0.03	23	359
C-312168	GLOC-7_38-40	0.3	7.7	39	17.6	9.12	0.03	23	378
C-312166	GLOC-7_40-42	0.4	7.3	41	12.4	8.89	0.03	22	361
C-312173	GLOC-7_42-44	0.3	8.0	42	16.0	9.35	0.03	24	417
C-312180	GLOC-7_44-46	0.3	8.5	45	13.5	9.55	0.03	25	441
C-312171	GLOC-7_46-48	0.4	8.8	42	17.7	9.75	0.03	26	437
C-312156	GLOC-7_48-50	0.3	8.5	43	15.5	9.02	0.03	25	477
C-312174	GLOC-7_50-52	0.4	9.6	43	17.0	9.95	0.03	27	508
C-312161	GLOC-7_52-54	0.2	10.2	44	16.2	9.82	0.04	26	527

Table 1–6. Core sediment sample analyses for GLOC-7, Grand Lake O' the Cherokees 4 kilometers downstream from Highway 2 bridge. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Ni ppm	P ppm	Pb ppm	Rb ppm	Sr ppm	V ppm	Zn ppm
C-312182	GLOC-7_0-2	18.1	540	28.1	73.9	93.8	63	336
C-312172	GLOC-7_2-4	18.3	520	29.2	77.9	90.0	63	362
C-312159	GLOC-7_4-6	19.5	530	27.7	80.1	94.4	68	231
C-312160	GLOC-7_6-8	23.8	840	53.9	95.0	97.6	85	463
C-312169	GLOC-7_8-10	24.4	760	36.8	97.0	98.4	84	339
C-312154	GLOC-7_10-12	20.5	580	21.9	84.0	99.3	72	160
C-312176	GLOC-7_12-14	19.4	510	25.1	85.0	99.4	70	211
C-312183	GLOC-7_14-16	24.8	610	24.9	90.3	98.7	77	166
C-312162	GLOC-7_17-18	19.4	460	20.1	80.3	98.3	67	152
C-312151	GLOC-7_18-20	24.6	550	22.3	89.1	108	80	190
C-312155	GLOC-7_20-22	22.5	590	32.5	86.0	97.0	76	315
C-312158	GLOC-7_22-24	20.7	560	24.1	77.0	95.8	66	275
C-312179	GLOC-7_24-26	25.1	680	34.9	95.0	102	81	542
C-312167	GLOC-7_26-28	25.2	620	35.9	92.5	103	82	554
C-312178	GLOC-7_28-30	22.1	500	24.3	87.5	84.8	74	231
C-312170	GLOC-7_30-32	16.8	310	19.6	70.3	62.5	59	101
C-312157	GLOC-7_32-34	16.1	300	17.8	64.0	57.3	58	75
C-312163	GLOC-7_34-36	17.2	310	18.7	69.9	58.1	61	87
C-312177	GLOC-7_36-38	16.9	300	19.2	69.8	56.5	60	84
C-312168	GLOC-7_38-40	18.1	320	19.6	68.5	57.6	60	74
C-312166	GLOC-7_40-42	17.8	290	17.5	65.5	50.8	59	67
C-312173	GLOC-7_42-44	18.2	300	18.3	69.8	57.1	63	67
C-312180	GLOC-7_44-46	20.2	300	19.0	71.0	56.6	63	69
C-312171	GLOC-7_46-48	19.5	290	18.7	69.4	53.8	64	67
C-312156	GLOC-7_48-50	19.9	290	18.9	66.7	52.6	64	70
C-312174	GLOC-7_50-52	19.5	280	20.4	71.4	53.2	67	64
C-312161	GLOC-7_52-54	20.3	270	20.0	71.3	52.4	68	64

Table 1–7. Core sediment sample analyses for GLOC-8, Grand Lake O' the Cherokees at confluence of Spring and Neosho Rivers. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.
[\(Click to see Excel file.\)](#)

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Depth in inches	Depth in cm	Total C %	Carbonate C %	Organic C %	Al %	Ca %	Fe %
C-310682	GLOC-8_1-3	2	5	1.1	0.02	1.08	4.95	0.47	2.27
C-310709	GLOC-8_3-5	4	10	1.15	0.01	1.14	5.32	0.49	2.41
C-310717	GLOC-8_5-7	6	15	1.13	0.01	1.12	5.2	0.46	2.37
C-310695	GLOC-8_7-9	8	20	1.61	<0.003	1.61	5.84	0.42	2.51
C-310692	GLOC-8_9-11	10	25	1.2	0.01	1.19	5.41	0.44	2.5
C-310697	GLOC-8_11-13	12	30	1.39	<0.003	1.39	5.82	0.44	2.69
C-310715	GLOC-8_13-15	14	35	1.38	0.01	1.37	5.26	0.39	2.34
C-310701	GLOC-8_15-17	16	40	1.17	0.01	1.16	3.89	0.3	1.6
C-310704	GLOC-8_17-19	18	45	1.25	0.01	1.24	3.81	0.29	1.54
C-310691	GLOC-8_19-21	20	50	1.27	0.03	1.24	3.95	0.32	1.71
C-310687	GLOC-8_21-23	22	55	1.13	0.01	1.12	4.19	0.32	1.75
C-310699	GLOC-8_23-25	24	60	1.34	0.03	1.31	5.13	0.67	2.5
C-310694	GLOC-8_25-27	26	65	0.86	0.01	0.85	4.27	0.33	1.72
C-310683	GLOC-8_27-29	28	70	0.74	<0.003	0.74	4.1	0.31	1.52
C-310689	GLOC-8_29-31	30	75	0.53	<0.003	0.53	3.97	0.29	1.53

Lab no.	Field no.	K %	Mg %	Na %	S %	Ti %	As ppm	Ba ppm	Bi ppm
C-310682	GLOC-8_1-3	1.29	0.36	0.46	0.04	0.23	7	453	0.25
C-310709	GLOC-8_3-5	1.35	0.4	0.47	0.04	0.24	7	468	0.27
C-310717	GLOC-8_5-7	1.31	0.38	0.44	0.04	0.24	6	473	0.28
C-310695	GLOC-8_7-9	1.35	0.38	0.38	0.04	0.28	7	509	0.38
C-310692	GLOC-8_9-11	1.21	0.39	0.4	0.04	0.24	7	476	0.31
C-310697	GLOC-8_11-13	1.41	0.43	0.42	0.04	0.26	7	491	0.33
C-310715	GLOC-8_13-15	1.31	0.35	0.42	0.04	0.26	6	482	0.3
C-310701	GLOC-8_15-17	1.18	0.21	0.42	0.03	0.23	4	474	0.18
C-310704	GLOC-8_17-19	1.14	0.22	0.4	0.03	0.22	4	458	0.19
C-310691	GLOC-8_19-21	1.16	0.22	0.41	0.03	0.23	5	473	0.19
C-310687	GLOC-8_21-23	1.13	0.23	0.4	0.04	0.23	6	506	0.2
C-310699	GLOC-8_23-25	1.12	0.4	0.24	0.09	0.25	9	409	0.29
C-310694	GLOC-8_25-27	1.2	0.25	0.42	0.02	0.24	4	483	0.19
C-310683	GLOC-8_27-29	1.2	0.24	0.44	0.03	0.24	4	528	0.19
C-310689	GLOC-8_29-31	1.2	0.23	0.43	0.01	0.21	3	487	0.19

Table 1–7. Core sediment sample analyses for GLOC-8, Grand Lake O' the Cherokees at confluence of Spring and Neosho Rivers. Analyses by ICP-AES/MS after four-acid digestion, except for forms of carbon.—Continued [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; cm, centimeter; %, percent; ppm, parts per million]

Lab no.	Field no.	Cd ppm	Co ppm	Cr ppm	Cu ppm	Ga ppm	In ppm	Li ppm	Mn ppm
C-310682	GLOC-8_1-3	20.5	11.4	47	21.4	11.7	0.07	30	552
C-310709	GLOC-8_3-5	16.4	11.5	52	23.3	12.1	0.07	32	530
C-310717	GLOC-8_5-7	19.5	11.1	50	25	11.8	0.07	32	521
C-310695	GLOC-8_7-9	25	13.6	58	32.8	13.4	0.07	36	647
C-310692	GLOC-8_9-11	21.9	12.5	67	28.1	12.8	0.07	33	522
C-310697	GLOC-8_11-13	22.1	12.8	51	26.7	13.5	0.05	36	583
C-310715	GLOC-8_13-15	16.9	12.2	48	26.1	12	0.05	32	611
C-310701	GLOC-8_15-17	3.5	9.8	42	17.8	8.3	0.03	24	428
C-310704	GLOC-8_17-19	5.8	8.3	29	16.8	8.7	0.03	24	445
C-310691	GLOC-8_19-21	3.1	10.1	44	15.3	8.4	0.03	24	893
C-310687	GLOC-8_21-23	1.9	11	37	13.6	9.1	0.04	26	399
C-310699	GLOC-8_23-25	2	11.7	49	21.2	12.3	0.04	33	556
C-310694	GLOC-8_25-27	1.1	10.1	34	14.4	9.4	0.03	26	731
C-310683	GLOC-8_27-29	0.9	8.7	36	14.5	9	0.03	25	513
C-310689	GLOC-8_29-31	0.7	9.7	33	13.2	9.4	0.03	23	546

Lab no.	Field no.	Ni ppm	P ppm	Pb ppm	Rb ppm	Sr ppm	V ppm	Zn ppm
C-310682	GLOC-8_1-3	23.8	740	82	77.2	74.9	76	1,460
C-310709	GLOC-8_3-5	25.8	740	85	85	76	81	1,530
C-310717	GLOC-8_5-7	24.5	760	102	82.6	74.1	79	1,580
C-310695	GLOC-8_7-9	27.9	780	204	88.3	73.9	89	2,030
C-310692	GLOC-8_9-11	31	750	115	80.1	74	87	1,570
C-310697	GLOC-8_11-13	28.8	640	112	87.4	77.5	88	1,770
C-310715	GLOC-8_13-15	27.9	600	130	85	74.1	83	1,720
C-310701	GLOC-8_15-17	24.3	570	62.4	69.6	66.6	56	608
C-310704	GLOC-8_17-19	17.9	580	70.2	71.7	63.9	59	969
C-310691	GLOC-8_19-21	19.3	540	51.1	68.9	66.7	58	456
C-310687	GLOC-8_21-23	18.6	410	37	72	68	63	277
C-310699	GLOC-8_23-25	24.4	760	34.9	82.5	72.3	83	478
C-310694	GLOC-8_25-27	18.5	360	21.6	73.4	67.3	61	120
C-310683	GLOC-8_27-29	19.3	370	22.3	72.5	68.7	60	118
C-310689	GLOC-8_29-31	16.1	380	19.4	72.9	64.6	49	92

Appendix 2. Data Quality Control and Quality Assessment

Sources of Geochemical Variation and Methods for Assuring Data Quality

There are several potential sources and levels of variation in geochemical data—for instance, between-site variation, within-site variation, sample inhomogeneity, and analytical variation. A complete analysis of variation (ANOVA) sampling design generally shows that the primary variation in geochemical data is that found between individual samples at different and widely spaced sites (between-site variation). This variation is due to differences in sample parent material, local geology, mineralization processes (Rose and others, 1979), and possible anthropogenic influences. This variation provides areal geochemical contrasts and is the basis upon which geochemical exploration programs operate.

The next level of geochemical variation is known as within-site variation. This variation can be reduced by compositing several subsamples within the immediate area of sample collection, resulting in a more representative sample of the entire site. Variation is also found within any single sample due to the heterogeneity of the sampled material. This sample variation can be reduced by good sample preparation procedures (Fey and others, 2009). A process of crushing, grinding, mixing, and splitting the sample typically creates a very fine homogenous powder from the original heterogeneous material. All of the sample preparation processes were performed by USGS personnel in either Oklahoma City or Denver. Laboratory analytical procedures also can be sources of variation in the geochemical data. These sources include differences in analysts, dissolution procedures, analytical instruments, instrument calibration errors, and instrument drift. The combined variation due to sample preparation, aliquot size, and analytical procedures (commonly called analytical variation) can be assessed by using standard reference materials, analytical duplicates, and procedure blanks (Dux, 1986).

Quality Control Samples

The SGS Minerals Services contract laboratory analyzed quality control samples along with the field samples. The SGS laboratories are ISO 17025 accredited, certified by the International Organization for Standardization (ISO). A standard reference material (SRM) was run for every 20 samples; this material was the Canadian soil standard CCRMP SO-3 (Canadian Certified Reference Materials Project: <http://www.nrcan.gc.ca/smm-mms/tect-tech/ccrmp/cer-cer/so-2-eng.htm>). Laboratory personnel also processed and analyzed about one out of every 20 samples twice as analytical duplicates. Procedure blanks, which are solutions of the acid reagents carried through the entire process, were also analyzed at a rate of one for every 20 samples. In this study, the data for these laboratory internal quality control samples were requested from the contract laboratory so that the analytical variation could be assessed.

In addition to the SGS internal laboratory quality-control samples, the USGS submitted additional quality control samples as blind samples. These included sample splits for analytical duplicates (for precision assessment), three National Institute of Standards and Technology (NIST) standards (NIST-2709, NIST-2711, and NIST-8704), and a suite of USGS-prepared standard reference materials (SAR-L, SAR-M, DGPM-1, and GSP-QC)¹ (for precision and accuracy assessment). These blind quality control samples were submitted to SGS Minerals Services at a rate of ten percent.

¹ SAR-L and SAR-M: sediment, Animas River watershed, Colo., low and moderate levels of contaminant constituents, respectively; DGPM-1: disseminated gold ore, Pinson Mine, Nev.; GSP-QC: granodiorite, Silver Plume, Colo.

Quality Assurance/Quality Control Measurement

A quality management system for a geochemical survey includes both quality assurance (QA) and quality control (QC) elements. The QA focus is mainly in the analytical laboratory environment. Under the QA umbrella, the components of standard operating procedures, instrument logs, training records, data acceptance/rejection criteria, and lab audits are covered (Dux, 1986). The QA element is not easily measured. However, the QC element provides measures of the accuracy and precision of geochemical data produced by an analytical method. The precision of an analytical method can be measured by the percent relative standard deviation (percent RSD) of data for a number of runs of a particular sample or standard; it is calculated by dividing the standard deviation (SD) by the mean and multiplying by 100. This most common version of the SD is defined as the square root of the quantity {"sum of squares of deviations of individual results from the mean, divided by one less than the number of results in the set"} (Dux, 1986):

$$SD = \sqrt{\sum_{i=1}^n (X_i - \bar{X})^2 / n - 1}$$

This precision was calculated for each element, for each SRM. The resulting eight precision values were then pooled together to derive an overall RSD for each element, using the following relation:

$$SD_{\text{pooled}} = \sqrt{(n_1 SD_1^2 + n_2 SD_2^2 + n_3 SD_3^2 + n_4 SD_4^2 + n_5 SD_5^2 + n_6 SD_6^2 + n_7 SD_7^2 + n_8 SD_8^2) / (n_1 + n_2 + n_3 + n_4 + n_5 + n_6 + n_7 + n_8)}$$

Where $n_{1...8}$ = number of pairs for a particular element, and $SD_{1...8}$ = individual standard deviation calculated for each SRM and element. The precision as represented by the percent RSD calculated from this method is shown in column 8 of table 2–1.

Table 2–1. Summary statistics for assessing analytical precision for carbonate carbon, total carbon, and 26 elements by ICP-AES/MS. Percent RSD, mean of relative percent differences, and pooled standard deviation are three separate measurements of analytical precision. ([Click to see Excel file.](#))

[ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; RPD, relative percent differences; RSD, relative standard deviation; SRMS, standard reference materials; %, percent; ppm, parts per million]

Element	Unit	Pairs (k)	Min	Max	Mean	Standard deviation based on duplicates	Percent RSD based on duplicate pairs	Mean of RPD between pairs	Percent RSD based on SRMS (pooled)
Carbonate									
C	%	22	0.003	1.47	0.03	0.0067	2.00	20.0	14.0
Total C	%	25	0.27	1.55	0.90	0.0179	1.99	2.21	4.56
Al	%	25	2.94	6.16	4.56	0.0780	1.71	1.90	2.61
Ca	%	25	0.18	0.79	0.37	0.0087	2.36	2.70	2.38
Fe	%	25	1.39	2.64	1.96	0.0352	1.79	2.02	2.88
K	%	25	0.77	1.57	1.26	0.0220	1.75	1.81	3.13
Mg	%	25	0.17	0.51	0.31	0.0071	2.25	2.39	2.65
Na	%	25	0.22	0.68	0.43	0.0088	2.03	2.29	3.59
S	%	23	0.01	0.06	0.04	0.0000	0.00	0.00	4.20
Ti	%	25	0.16	0.26	0.22	0.0132	6.12	6.64	5.18
As	ppm	25	3.0	9.0	5.59	0.616	11.0	7.04	9.20
Ba	ppm	25	299	527	432	8.71	2.02	2.18	1.88
Bi	ppm	20	0.15	0.39	0.22	0.010	4.46	5.60	15.5
Cd	ppm	25	0.2	24.4	4.00	0.317	7.92	9.78	19.0
Co	ppm	25	7.0	14.3	10.0	0.370	3.69	3.82	11.6
Cr	ppm	25	32	59	45.2	2.63	5.83	5.46	7.66
Cu	ppm	25	12.3	40	18.3	0.932	5.09	5.52	6.96
Ga	ppm	25	6.92	14.6	10.6	0.283	2.68	2.98	8.77
In	ppm	25	0.03	0.07	0.04	0.003	8.03	3.69	13.5
Li	ppm	25	19	36	26.3	0.693	2.63	3.08	3.53
Mn	ppm	25	323	980	543	15.5	2.86	2.83	4.39
Ni	ppm	25	14	32.6	22.0	1.26	5.70	5.19	9.94
P	ppm	25	230	930	503	12.6	2.52	2.45	2.69
Pb	ppm	25	15	248	52.7	2.58	4.89	3.99	6.00
Rb	ppm	25	50.1	100	75.1	2.18	2.91	2.80	6.52
Sr	ppm	25	37.4	116	74.4	1.43	1.93	2.39	1.92
V	ppm	25	46	91	67.4	1.21	1.79	1.59	2.64
Zn	ppm	25	48	3,500	586	32.0	5.47	3.10	3.67

A second way to assess the precision of a technique is to calculate the average of the relative percent differences (RPDs) between pairs of duplicate measurements: samples that were split in the laboratory after field collection and then processed and analyzed as two separate samples. In this study, there were 25 pairs of duplicate measurements for most elements. The version of precision thus calculated is:

$$\text{Precision} = \{\text{Average of } \sum (\text{absolute value } (X_1 - X_2) / (X_1 + X_2)) / 2\} * 100$$

where X_1 and X_2 are the measurement values of each duplicate pair. This precision is shown in column 9 of table 2–1.

Another way to estimate precision from analytical duplicate pairs is to calculate the standard deviation as the square root of the quantity {sum of squares of the difference between the duplicate results (R), divided by two times the number of sets of duplicate samples} (Dux, 1986).

$$SD_{dupe} = \sqrt{\sum R^2 / 2k}$$

The percent RSD is then again calculated by dividing the standard deviation (SD_{dupe}) by the mean and multiplying by 100. This third estimate for precision in this study is shown in column 10 of table 2–1.

The accuracy of an analytical method is measured by the percent recovery, which is calculated by dividing the mean concentration of replicate analyses by the target value of the standard reference material used and multiplying by 100. Target values for standard reference materials may consist of certified values or, when certified values are not available, informational values. The percent recovery derived from informational values is not as reliable as the percent recovery calculated from certified values, but it is still useful. In general, analytical determinations become less accurate and precise as data values approach the lower or upper reporting limits. Measurements of precision and accuracy are best when elemental concentrations fall within the middle of the determination range for a specified analytical method and element. Percent RSD and percent recovery values are more robust when calculated on mean values greater than five times the method's lower reporting limit.

Possible contamination during the analytical procedure is assessed through the use of procedure blanks. Blanks are defined based on the sample medium and analytical method and are processed concurrently with samples to determine whether contamination has occurred during the sample dissolution stages or whether cross-sample contamination has occurred in the analytical instrument during a sample run. Commonly, blanks are a set of all the reagents used in the sample processing and analysis procedure. Procedure blanks were prepared, analyzed, and results provided by SGS Minerals to the USGS. The blank results are shown in table 1–1; they show no contamination effects from reagents or the analytical procedures.

Quality Assessment of Data

Eight different reference materials were analyzed with the samples for the ICP-AES/MS method. Five of these materials have certified values (NIST 2709, NIST 2711, NIST 8704, USGS DGPM-1, CCRMP SO-3), and three are in-house USGS materials that have been submitted to the contract laboratory over a period of years but do not have certified values (USGS SAR-L, USGS SAR-M, USGS GSP-QC). No single certified reference material covers all of the elements analyzed; the use of several different certified SRMs allowed for all elements except bismuth (Bi) and lithium (Li) to have certified accuracy (percent recovery) assessments calculated. Comparison of the concentration values for bismuth

and lithium for USGS SAR-L, SAR-M, and GSP-QC with their respective historical values allowed for noncertified percent recovery calculations to be made for those two elements.

Precision Analysis

As outlined above, precision was calculated using three different approaches. Analysis of 25 duplicate pairs yielded a percent RSD and a mean of relative percent differences for all elements. Multiple analyses of each reference material allowed for a calculation of a percent RSD for each. These RSDs were then pooled to produce a third overall measure of precision for the dataset. The three calculated precision values are plotted in figure 2–1. In general, the precision values mostly plot at better than <10 percent by all three methods. Tables 2–2 through 2–9 include columns for the standard deviation and percent relative standard deviation for each respective reference material. These are the values used in the pooled standard deviation calculation discussed above. A brief description follows for those elements whose RSD calculations were above ten percent for one or more of the precision parameters: carbonate carbon, arsenic, bismuth, cadmium, cobalt, indium, and nickel.

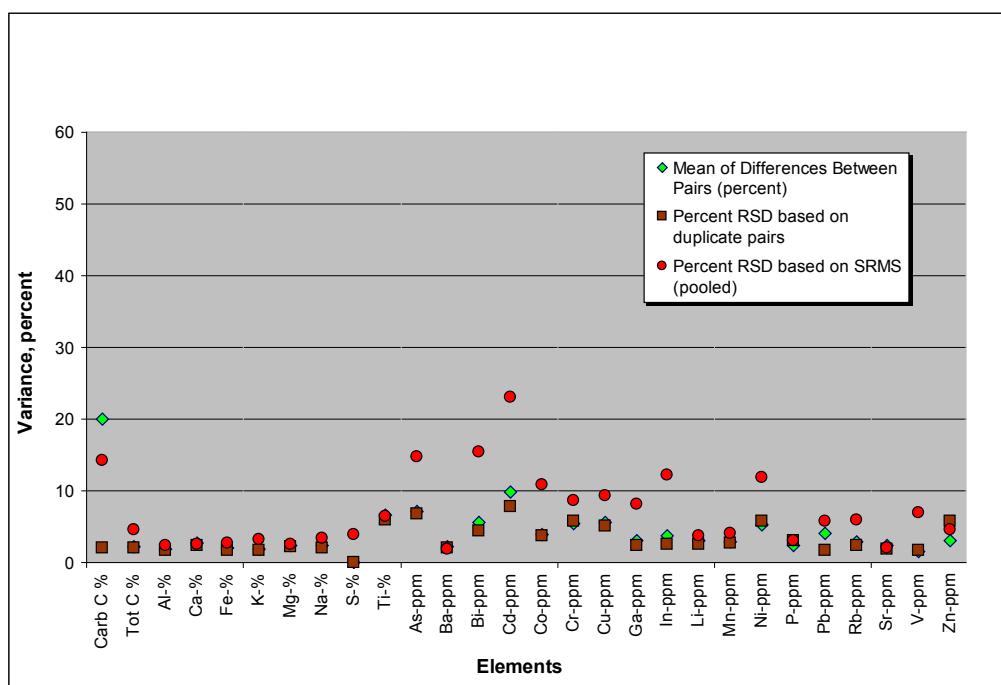


Figure 2–1. Precision plot for ICP-AES/MS and carbon analyses based on three estimates of variance. [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; %, percent; ppm, parts per million; RSD, relative standard deviation; SRMS, standard reference materials]

The high variability for carbonate carbon based on the mean of differences between pairs, 20 percent, is related to the weight contributed to the calculation from samples whose carbonate carbon contents are very low, near the detection limit. For example, the percent difference between duplicate analyses of 0.01 and 0.02 is 67 percent, whereas the percent difference between duplicate analyses of 1.15 and 1.13 is only 1.7 percent. Thus, the sample pairs with very low carbonate carbon tend to skew this calculation of the precision parameter.

The pooled RSD for arsenic was 14.7 percent, but this value was skewed by the individual RSD for the reference material CCRMP SO-3, which was 29 percent. Discarding this one value resulted in a pooled RSD for arsenic of 8 percent.

The precision for bismuth based on the duplicate pairs and the mean of the RPDs is about 5 percent (table 2–1). The pooled RSD based on the eight reference materials is 15.5 percent. This greater value is influenced by those standards that had bismuth concentrations at less than 1 ppm. At this level, distributions of trace elements are much less homogeneous. The same situation and explanation applies to cadmium.

The pooled estimate for RSD for cobalt was 10.8 percent. A single outlier value for the SRM NIST 8704 of 5.1 ppm versus a target of 13.6 ppm resulted in this high RSD. Eliminating that outlier value results in a pooled RSD estimate for cobalt of 4.7 percent.

A single outlier value for the SRM NIST 8704 also influenced the pooled RSD estimate for indium. With the outlier the estimate is 12.2 percent. Removing the outlier (0.03 ppm versus a mean of 0.08 ppm) results in a pooled RSD estimate of 5.1 percent.

Accuracy Analysis

SRM NIST 2709

The SRM NIST 2709 was analyzed six times along with the samples. Table 2–2 gives the percent RSD and percent recovery results for this standard, and figure 2–2 shows graphically the accuracy plot. Those elements that show “n/a” in table 2–2 or have no symbol plotted on figure 2–2 do not have a certified value for comparison. All plotted elements except chromium and nickel fall within ± 15 percent of the target values. The chromium recovery is low, 76 percent, due to the difficulty of the four-acid digestion to dissolve the refractory mineral phase in which chromium resides.

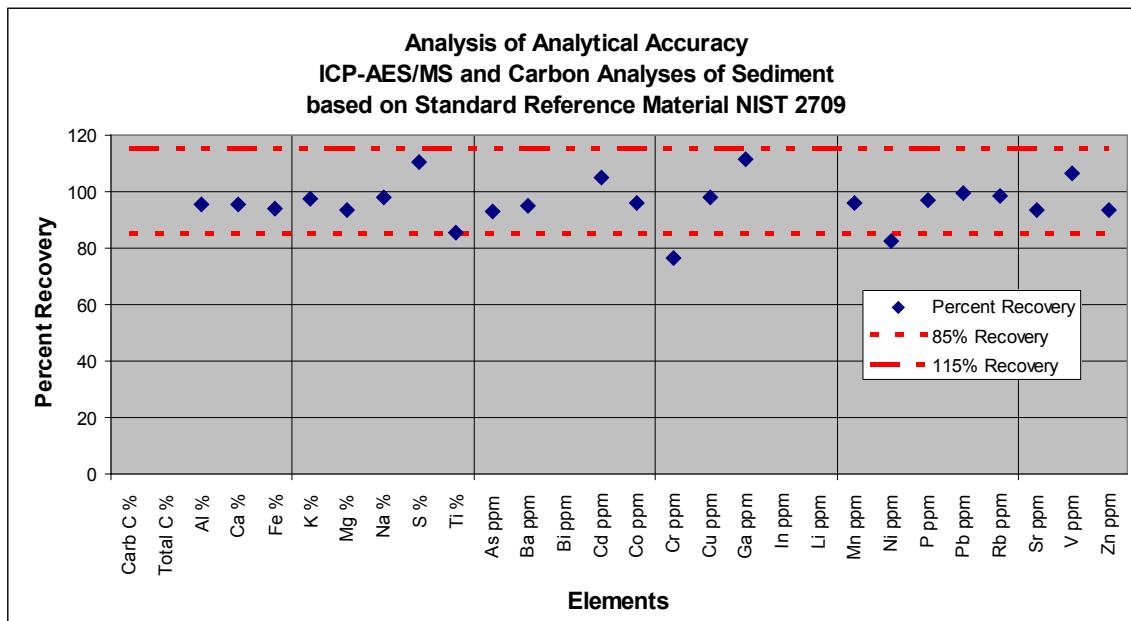


Figure 2–2. Accuracy plot for ICP-AES/MS and carbon analyses for standard reference material NIST 2709. [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; NIST, National Institute of Standards and Technology; %, percent; ppm, parts per million]

Table 2–2. Summary statistics for assessing analytical variation and accuracy for the standard reference material NIST 2709 determined after a four-acid total digestion of sediment samples by the ICP-AES-MS package at SGS Minerals and for total and carbonate carbon. ([Click to see Excel file.](#))

[NIST, National Institute of Standards and Technology; ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; RL, reporting limit; n, number of analyses of NIST 2709; RSD, relative standard deviation; %, percent; ppm, parts per million; n/a, not applicable]

Element	Units	RL	n	Target value	Mean	Standard deviation	% RSD	% Recovery
Al	%	0.01	6	7.50	7.16	0.12	1.74	96
Ca	%	0.01	6	1.89	1.81	0.01	0.81	96
Fe	%	0.01	6	3.50	3.29	0.07	2.25	94
K	%	0.01	6	2.03	1.98	0.04	2.24	97
Mg	%	0.01	6	1.51	1.41	0.03	1.88	94
Na	%	0.01	6	1.16	1.14	0.01	1.29	98
S	%	0.01	6	0.062	0.10	0.00	4.15	110
Ti	%	0.01	6	0.34	0.29	0.01	2.58	85
As	ppm	1	6	17.7	16.5	0.84	5.07	93
Ba	ppm	5	6	968	921	14.1	1.5	95
Bi	ppm	0.04	6	n/a	0.29	0.02	5.32	n/a
Cd	ppm	0.1	6	0.38	0.40	0.00	0.00	105
Co	ppm	0.1	6	13.4	12.8	0.38	2.9	96
Cr	ppm	1	6	130	99.3	3.50	3.53	76
Cu	ppm	0.5	6	34.6	33.8	2.42	7.2	98
Ga	ppm	0.05	6	14	15.6	0.57	3.7	111
In	ppm	0.02	6	n/a	0.05	0.00	7.9	n/a
Li	ppm	1	6	n/a	51	1.41	2.8	n/a
Mn	ppm	5	6	538	516	13.3	2.6	96
Ni	ppm	0.5	6	88.0	72.5	1.86	2.6	82
P	ppm	50	6	0.062	602	29.3	4.9	99
Pb	ppm	0.5	6	18.9	18.8	1.67	8.9	100
Rb	ppm	0.2	6	n/a	94.4	3.16	3.3	n/a
Sr	ppm	0.5	6	231	216	2.07	1.0	94
V	ppm	1	6	112	120	2.51	2.1	107
Zn	ppm	1	6	106	99.3	3.50	3.5	94
Total C	%	0.01	6	n/a	1.16	0.03	2.8	n/a
Carbonate C	%	0.003	6	n/a	0.25	0.00	1.62	n/a

SRM NIST 2711

The SRM NIST 2711 was analyzed six times along with the samples. Table 2–3 gives the percent RSD and percent recovery results for this standard, and figure 2–3 shows graphically the accuracy plot. Those elements that show “n/a” in table 2–3 or have no symbol plotted on figure 2–3 do not have a certified value for comparison. All plotted elements except chromium and nickel fall within ± 15 percent of the target values. The chromium recovery is low, 79 percent, due to the difficulty of the four-acid digestion to dissolve the refractory mineral phase in which chromium resides.

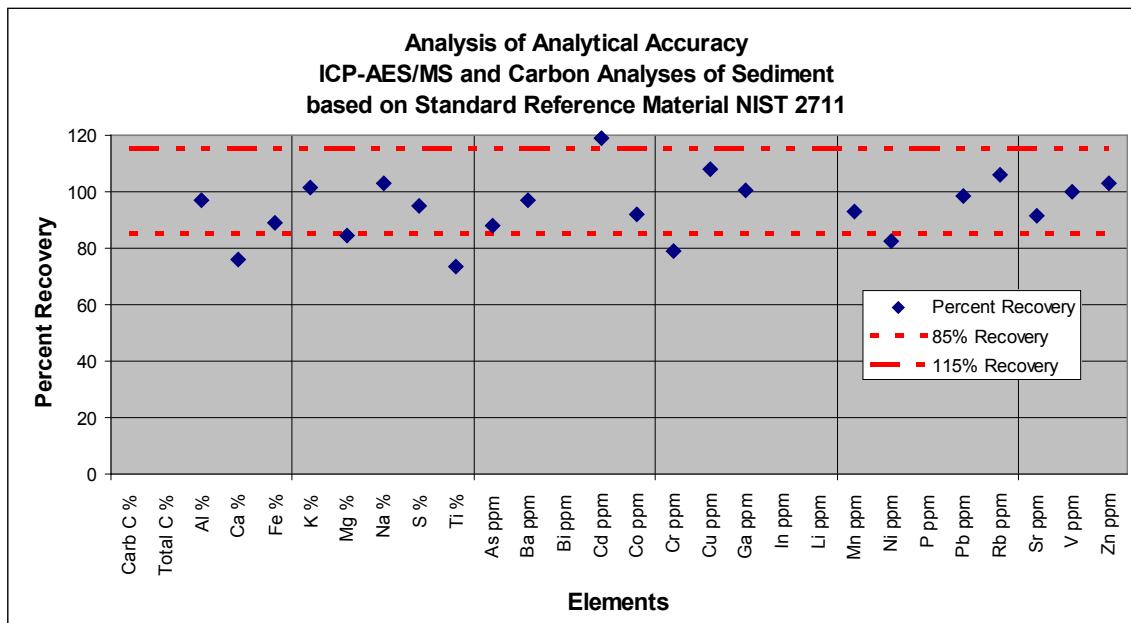


Figure 2–3. Accuracy plot for ICP-AES/MS and carbon analyses for standard reference material NIST 2711. [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; NIST, National Institute of Standards and Technology; %, percent; ppm, parts per million]

Table 2–3. Summary statistics for assessing analytical variation and accuracy for the standard reference material NIST 2711 determined after a four-acid total digestion of sediment samples by the ICP-AES/MS package at SGS Minerals and for total and carbonate carbon. ([Click to see Excel file.](#))

[NIST, National Institute of Standards and Technology; ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; RL, reporting limit ; n, number of analyses of NIST 2711; RSD, relative standard deviation; %, percent; ppm, parts per million; n/a, not applicable]

Element	Units	RL	n	Target value	Mean	Standard deviation	% RSD	% Recovery
Al	%	0.01	6	6.53	6.34	0.18	2.87	97
Ca	%	0.01	6	2.88	2.19	0.06	2.64	76
Fe	%	0.01	6	2.89	2.57	0.06	2.43	89
K	%	0.01	6	2.45	2.49	0.08	3.13	102
Mg	%	0.01	6	1.05	0.89	0.03	3.26	84
Na	%	0.01	6	1.14	1.17	0.03	2.61	103
S	%	0.01	6	0.086	0.04	0.00	0.00	46
Ti	%	0.01	6	0.306	0.23	0.01	3.72	74
As	ppm	1	6	105	92.3	1.21	1.31	88
Ba	ppm	5	6	726	703	7.52	1.07	97
Bi	ppm	0.04	6	n/a	2.72	0.04	1.46	n/a
Cd	ppm	0.1	6	41.7	49.7	2.51	5.06	119
Co	ppm	0.1	6	10	9.18	0.32	3.47	92
Cr	ppm	1	6	47	37.2	2.71	7.30	79
Cu	ppm	0.5	6	114	123	5.38	4.37	108
Ga	ppm	0.05	6	15	15.1	0.79	5.25	101
In	ppm	0.02	6	n/a	1.13	0.04	3.90	n/a
Li	ppm	1	6	n/a	24.2	0.75	3.11	n/a
Mn	ppm	5	6	638	592	16.4	2.77	93
Ni	ppm	0.5	6	20.6	17.0	0.60	3.56	82
P	ppm	50	6	860	793	10.3	1.30	92
Pb	ppm	0.5	6	1,162	1,142	61.8	5.41	98
Rb	ppm	0.2	6	n/a	116	4.63	3.98	n/a
Sr	ppm	0.5	6	245	225	3.27	1.46	92
V	ppm	1	6	81.6	81.5	3.02	3.70	100
Zn	ppm	1	6	350	361	12.3	3.42	103
Total C	%	0.01	6	n/a	1.55	0.06	3.81	n/a
Carbonate C	%	0.003	6	n/a	0.34	0.01	1.59	n/a

SRM NIST 8704

The SRM NIST 8704 was analyzed six times along with the samples. Table 2–4 gives the percent RSD and percent recovery results for this standard, and figure 2–4 shows graphically the accuracy plot. Those elements that show “n/a” in table 2–4 or have no symbol plotted on figure 2–4 do not have a certified value for comparison. All plotted elements except titanium, arsenic, and chromium fall within ± 15 percent of the target values. The recovery for titanium was only 43 percent, and the recovery for chromium was 81 percent; both of these elements can occur in mineral phases that are difficult to fully dissolve by the four-acid digestion procedure. The arsenic recovery, 74 percent, is related to the low precision and accuracy at low analytical levels (arsenic is present at less than 20 times the method reporting limit).

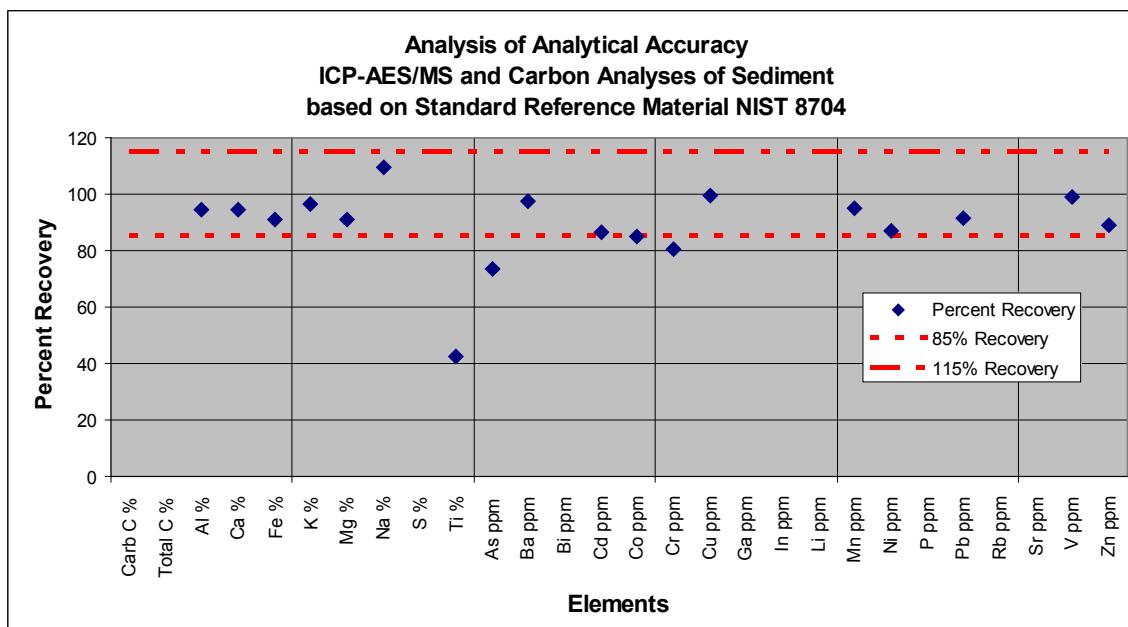


Figure 2–4. Accuracy plot for ICP-AES/MS and carbon analyses for standard reference material NIST 8704. [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; NIST, National Institute of Standards and Technology; %, percent; ppm, parts per million]

Table 2–4. Summary statistics for assessing analytical variation and accuracy for the standard reference material NIST 8704 determined after a four-acid total digestion of sediment samples by the ICP-AES/MS package at SGS Minerals and for total and carbonate carbon. ([Click to see Excel file.](#))

[NIST, National Institute of Standards and Technology; ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; RL, reporting limit; n, number of analyses of NIST 8704; RSD, relative standard deviation; %, percent; ppm, parts per million; n/a, not applicable]

Element	Units	RL	n	Target value	Mean	Standard deviation	% RSD	% Recovery
Al	%	0.01	6	6.1	5.77	0.09	1.6	95
Ca	%	0.01	6	2.64	2.49	0.05	2.0	94
Fe	%	0.01	6	3.97	3.61	0.07	1.9	91
K	%	0.01	6	2	1.93	0.04	2.2	97
Mg	%	0.01	6	1.20	1.09	0.02	2.1	91
Na	%	0.01	6	0.553	0.61	0.01	1.7	109
S	%	0.01	6	n/a	0.37	0.02	4.3	n/a
Ti	%	0.01	6	0.457	0.20	0.02	7.8	43
As	ppm	1	6	17.0	12.5	2.74	21.9	74
Ba	ppm	5	6	413	403	9.67	2.4	97
Bi	ppm	0.04	6	n/a	1.92	0.53	27.7	n/a
Cd	ppm	0.1	6	2.94	2.98	0.26	8.7	101
Co	ppm	0.1	6	13.6	2.55	1.08	42.3	85
Cr	ppm	1	6	122	11.6	3.18	27.5	81
Cu	ppm	0.5	6	87	98.5	2.51	2.5	100
Ga	ppm	0.05	6	n/a	86.8	5.88	6.8	n/a
In	ppm	0.02	6	n/a	13.0	2.53	19.5	n/a
Li	ppm	1	6	n/a	0.08	0.03	31.9	n/a
Mn	ppm	5	6	544	41.7	0.82	2.0	95
Ni	ppm	0.5	6	42.9	517	11.3	2.2	87
P	ppm	50	6	n/a	37.3	1.82	4.9	n/a
Pb	ppm	0.5	6	150	923	25.8	2.8	91
Rb	ppm	0.2	6	n/a	137	9.19	6.7	n/a
Sr	ppm	0.5	6	n/a	92.9	11.2	12.0	n/a
V	ppm	1	6	94.6	127	1.76	1.4	99
Zn	ppm	1	6	408	93.8	1.83	2.0	89
Total C	%	0.01	0	n/a	n/a	n/a	n/a	n/a
Carbonate C	%	0.003	0	n/a	n/a	n/a	n/a	n/a

SRM CCRMP SO-3

The SRM CCRMP SO-3 was analyzed eight times along with the samples. Table 2–5 gives the percent RSD and percent recovery results for this standard, and figure 2–5 shows graphically the accuracy plot. Those elements that show “n/a” in table 2–5 or have no symbol plotted on figure 2–5 do not have a certified value for comparison. All plotted elements except titanium, copper, and nickel fall within ± 15 percent of the target values. The titanium recovery was 80 percent, probably due to difficulties in obtaining a complete decomposition of refractory titanium phases using the four-acid digestion. The low accuracy estimates for copper and nickel (82 and 72 percent, respectively) may be due to those elements’ concentrations being near instrumental reporting limits.

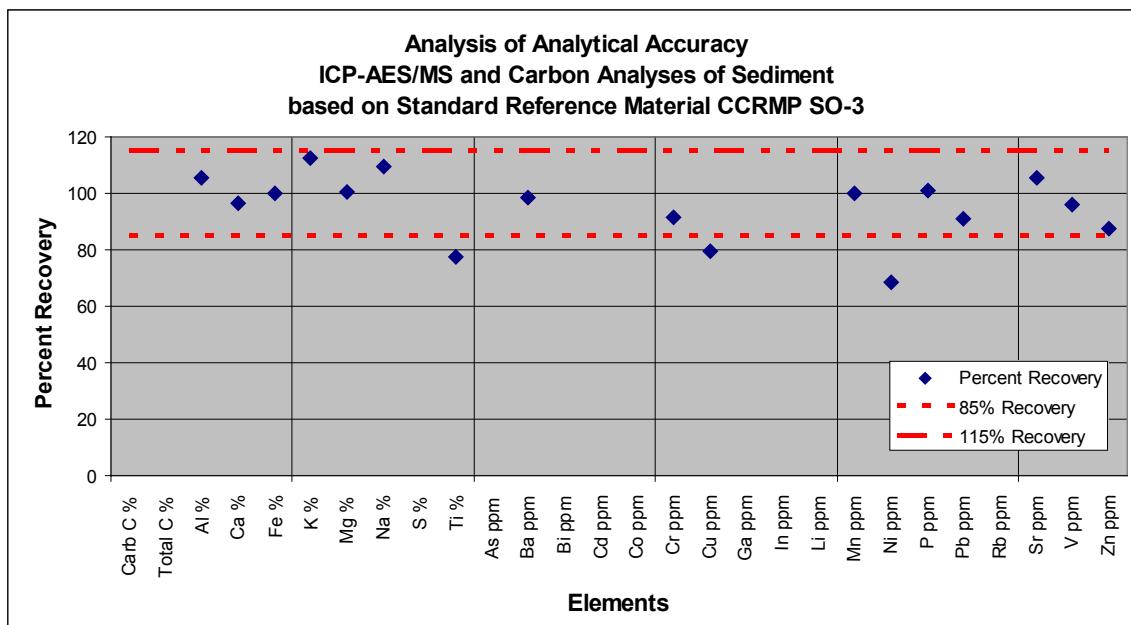


Figure 2–5. Accuracy plot for ICP-AES/MS and carbon analyses for standard reference material CCRMP SO-3. [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; CCRMP, Canadian Certified Reference Materials Project; %, percent; ppm, parts per million]

Table 2–5. Summary statistics for assessing analytical variation and accuracy for the standard reference material CCRMP SO-3 determined after a four-acid total digestion of sediment samples by the ICP-AES/MS package at SGS Minerals and for total and carbonate carbon. ([Click to see Excel file.](#))

[CCRMP, Canadian Certified Reference Materials Project; ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; RL, reporting limit; n, number of analyses of CCRMP SO-3; RSD, relative standard deviation; %, percent; ppm, parts per million; n/a, not applicable]

Element	Units	RL	n	Target value	Mean	Standard deviation	% RSD	% Recovery
Al	%	0.01	8	3.06	3.21	0.06	1.9	105
Ca	%	0.01	8	14.6	14.1	0.32	2.29	97
Fe	%	0.01	8	1.51	1.51	0.02	1.12	100
K	%	0.01	8	1.16	1.29	0.05	4.14	111
Mg	%	0.01	8	4.98	5.00	0.06	1.19	100
Na	%	0.01	8	0.74	0.80	0.02	3.02	108
S	%	0.01	8	n/a	0.02	0.00	0.00	n/a
Ti	%	0.01	8	0.2	0.16	0.02	9.88	80
As	ppm	1	8	n/a	2.00	0.58	28.9	n/a
Ba	ppm	5	8	296	292	5.46	1.87	99
Bi	ppm	0.04	8	n/a	0.08	0.01	15.5	n/a
Cd	ppm	0.1	8	n/a	0.14	0.05	37.6	n/a
Co	ppm	0.1	8	n/a	5.31	0.30	5.64	n/a
Cr	ppm	1	8	26	24.0	2.96	12.3	92
Cu	ppm	0.5	8	17	13.9	2.29	16.5	82
Ga	ppm	0.05	8	n/a	6.90	0.35	5.12	n/a
In	ppm	0.02	8	n/a	0.03	0.00	0.00	n/a
Li	ppm	1	8	n/a	12.6	0.52	4.10	n/a
Mn	ppm	5	8	520	520	5.87	1.13	100
Ni	ppm	0.5	8	16	11.5	1.87	16.3	72
P	ppm	50	8	480	484	11.3	2.33	101
Pb	ppm	0.5	8	14	12.9	0.62	4.85	92
Rb	ppm	0.2	8	n/a	37.6	1.40	3.72	n/a
Sr	ppm	0.5	8	217	227	4.66	2.05	105
V	ppm	1	8	38	36.7	0.71	1.93	96
Zn	ppm	1	8	52	46.1	3.41	7.39	89
Total C	%	0.01	8	n/a	6.66	0.11	1.60	n/a
Carbonate C	%	0.003	0	n/a	n/a	n/a	n/a	n/a

SRM USGS DGPM-1

The SRM USGS DGPM-1 was analyzed four times along with the samples. Table 2–6 gives the percent RSD and percent recovery results for this standard, and figure 2–6 shows graphically the accuracy plot. Those elements that show “n/a” in table 2–6 or have no symbol plotted on figure 2–6 do not have a certified value for comparison. All plotted elements except cadmium fall within ± 15 percent of the target values. The cadmium recovery, 83 percent, is quite acceptable given that the concentration is only three times the analytical reporting limit.

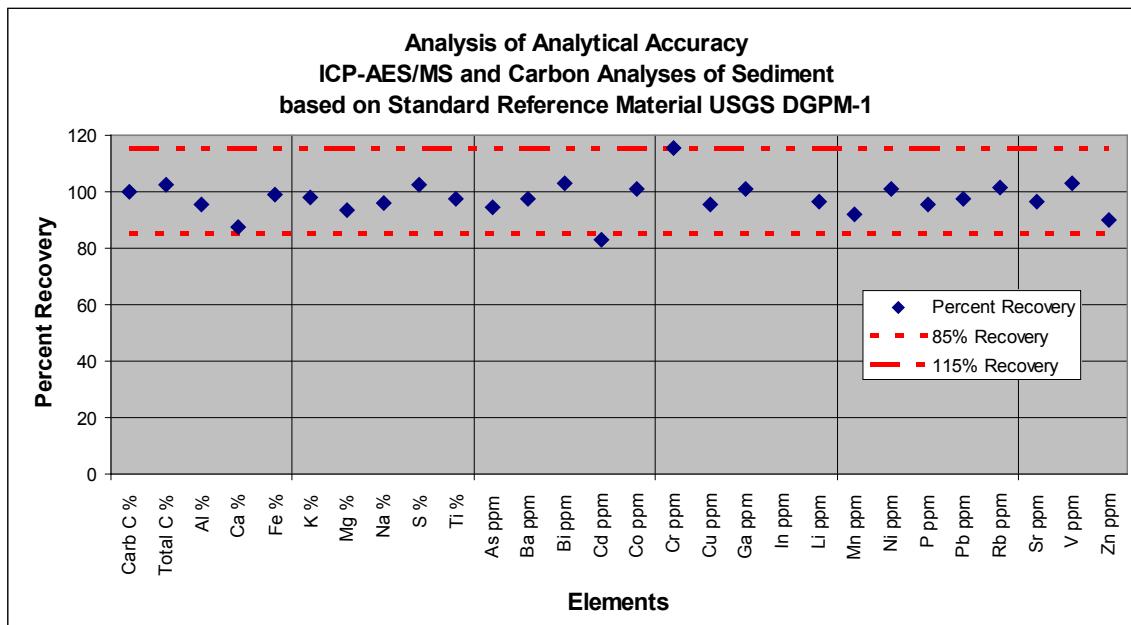


Figure 2–6. Accuracy plot for ICP-AES/MS and carbon analyses for standard reference material USGS DGPM-1. [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; USGS, U.S. Geological Survey; DGPM, disseminated gold ore, Pinson Mine, Nev.; %, percent; ppm, parts per million]

Table 2–6. Summary statistics for assessing analytical variation and accuracy for the standard reference material USGS DGPM-1 determined after a four-acid total digestion of sediment samples by the ICP-AES/MS package at SGS Minerals and for total and carbonate carbon. ([Click to see Excel file.](#))

[USGS, U.S. Geological Survey; DGPM, disseminated gold ore, Pinson Mine, Nev.; ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; RL, reporting limit; n, number of analyses of DGPM-1; RSD, relative standard deviation; %, percent; ppm, parts per million; n/a, not applicable]

Element	Units	RL	n	Target value	Mean	Standard deviation	% RSD	% Recovery
Al	%	0.01	4	5.06	4.88	0.17	3.4	96
Ca	%	0.01	4	0.157	0.14	0.01	3.54	90
Fe	%	0.01	4	1.34	1.33	0.04	3.04	99
K	%	0.01	4	2.27	2.24	0.07	2.99	99
Mg	%	0.01	4	0.337	0.32	0.01	1.81	95
Na	%	0.01	4	0.06	0.06	0.01	8.70	96
S	%	0.01	4	0.363	0.37	0.01	2.57	103
Ti	%	0.01	4	0.266	0.26	0.01	3.14	98
As	ppm	1	4	180	172	1.29	0.75	96
Ba	ppm	5	4	1,326	1,293	20.6	1.60	97
Bi	ppm	0.04	4	0.114	0.12	0.02	18.9	103
Cd	ppm	0.1	4	0.33	0.28	0.05	18.2	83
Co	ppm	0.1	4	1.36	1.38	0.15	10.9	101
Cr	ppm	1	4	97	112	17.05	15.2	115
Cu	ppm	0.5	4	13.7	13.1	1.71	13.0	95
Ga	ppm	0.05	4	10.8	10.9	0.46	4.25	101
In	ppm	0.02	4	n/a	<0.02	n/a	n/a	n/a
Li	ppm	1	4	39.6	38.2	0.96	2.50	97
Mn	ppm	5	4	28	25.7	2.87	11.2	92
Ni	ppm	0.5	4	11.4	11.5	3.56	31.0	101
P	ppm	50	4	418	400	8.16	2.04	96
Pb	ppm	0.5	4	9.8	9.58	0.66	6.84	98
Rb	ppm	0.2	4	90	91.6	4.77	5.21	102
Sr	ppm	0.5	4	91.5	88.2	1.43	1.63	96
V	ppm	1	4	106	109	2.16	1.98	103
Zn	ppm	1	4	24.4	22.0	1.15	5.25	90
Total C	%	0.01	4	0.10	0.10	0.01	9.34	103
Carbonate C	%	0.003	3	0.01	0.01	0.00	0.0	100

Reference Material USGS GSP-QC

In contrast to the first five SRMs discussed above, the reference material USGS GSP-QC is not certified nor used outside of the U.S. Geological Survey. It is a material similar to the certified SRM GSP-1 (Silver Plume Granite) but was developed at a later time, and so it is slightly different in elemental concentrations. This material was developed in-house at the Denver USGS laboratories for inclusion with samples sent to the SGS Minerals laboratory. During the period of September 2004 to November 2008, this material was analyzed 360 times by the contract laboratory for the ICP-AES/MS package as blind samples and 120 times for carbon. The results were compiled, and the means derived were used as target values for the accuracy assessments of this material submitted in this report. As a consequence, target values are available for all elements in the ICP-AES/MS package and for total and carbonate carbon.

The noncertified USGS reference material GSP-QC was analyzed five times along with the study samples. Table 2-7 gives the percent RSD and percent recovery results for this standard, and figure 2-7 shows graphically the accuracy plot. The only element that plotted outside of the 15-percent window was nickel at 82 percent, probably due to its low concentration.

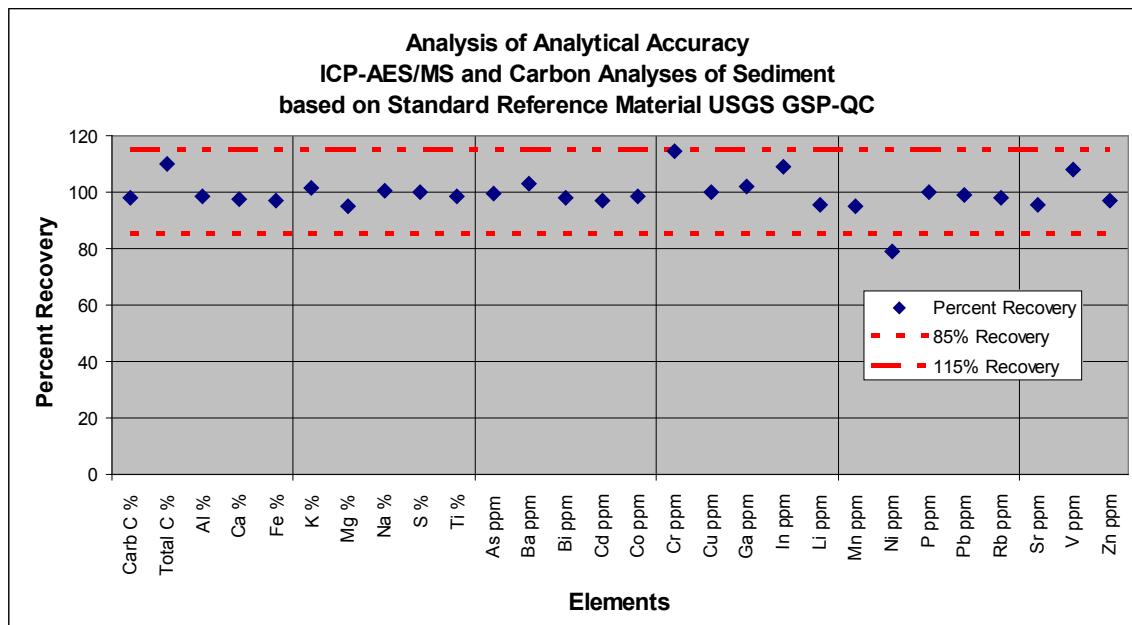


Figure 2-7. Accuracy plot for ICP-AES/MS and carbon analyses for reference material USGS GSP-QC. [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; USGS, U.S. Geological Survey; GSP-QC, granodiorite, Silver Plume, Colo.; %, percent; ppm, parts per million]

Table 2–7. Summary statistics for assessing analytical variation and accuracy for the reference material USGS GSP-QC determined after a four-acid total digestion of sediment samples by the ICP-AES/MS package at SGS Minerals and for total and carbonate carbon. ([Click to see Excel file.](#))

[USGS; U.S. Geological Survey; GSP-QC, granodiorite, Silver Plume, Colo.; ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; RL, reporting limit; n, number of analyses of GSP-QC; RSD, relative standard deviation; %, percent; ppm, parts per million; n/a, not applicable]

Element	Units	RL	n	Target value	Mean	Standard deviation	% RSD	% Recovery
Al	%	0.01	5	7.57	7.48	0.27	3.58	99
Ca	%	0.01	5	1.5	1.47	0.05	3.55	98
Fe	%	0.01	5	2.77	2.70	0.13	4.84	97
K	%	0.01	5	4.25	4.30	0.20	4.57	101
Mg	%	0.01	5	0.615	0.59	0.02	3.81	96
Na	%	0.01	5	1.87	1.88	0.08	4.00	101
S	%	0.01	5	0.074	0.07	0.00	6.62	100
Ti	%	0.01	5	0.339	0.33	0.02	6.46	99
As	ppm	1	5	31.4	31.2	2.04	6.54	99
Ba	ppm	5	5	1,310	1,343	37.8	2.81	103
Bi	ppm	0.04	5	4.28	4.21	0.17	4.12	98
Cd	ppm	0.1	5	0.227	0.22	0.04	18.1	97
Co	ppm	0.1	5	6.3	6.22	0.18	2.95	99
Cr	ppm	1	5	16.6	18.6	2.48	13.3	112
Cu	ppm	0.5	5	31.3	31.3	2.52	8.06	100
Ga	ppm	0.05	5	22.2	22.6	0.84	3.74	102
In	ppm	0.02	5	0.044	0.05	0.00	9.13	108
Li	ppm	1	5	34	32.7	1.51	4.61	96
Mn	ppm	5	5	281	270	10.3	3.81	96
Ni	ppm	0.5	5	11.5	9.48	1.05	11.1	82
P	ppm	50	5	1,230	1,228	29.3	2.38	100
Pb	ppm	0.5	5	40.4	40.1	1.32	3.29	99
Rb	ppm	0.2	5	228	224	14.9	6.66	98
Sr	ppm	0.5	5	226	218	8.59	3.95	96
V	ppm	1	5	73.3	78.1	3.55	4.55	106
Zn	ppm	1	5	117	114	4.82	4.23	97
Total C	%	0.01	5	n/a	0.20	0.00	2.26	n/a
Carbonate C	%	0.003	5	n/a	0.32	0.01	3.44	n/a

Reference Material USGS SAR-L

Like USGS GSP-QC, the reference material SAR-L is not certified nor used outside of the U.S. Geological Survey. It is a composite of contaminated sediment from the Animas River watershed in Colorado with low levels of contaminant constituents. This material was developed in-house at the Denver USGS laboratories for inclusion with samples sent to the SGS laboratory. During the period of September 2004 to November 2008, this material was analyzed 360 times by the SGS laboratory for the ICP-AES/MS package as blind samples and 150 times for carbon. The results were compiled, and the means derived were used as target values for the accuracy assessments of this material submitted in this report. As a consequence, target values are available for all elements in the ICP-AES/MS package and for total and carbonate carbon.

The noncertified USGS reference material SAR-L was analyzed seven times along with the study samples. Table 2–8 gives the percent RSD and percent recovery results for this standard, and figure 2–8 shows graphically the accuracy plot. All of the elements plot within ± 15 percent for percent recovery.

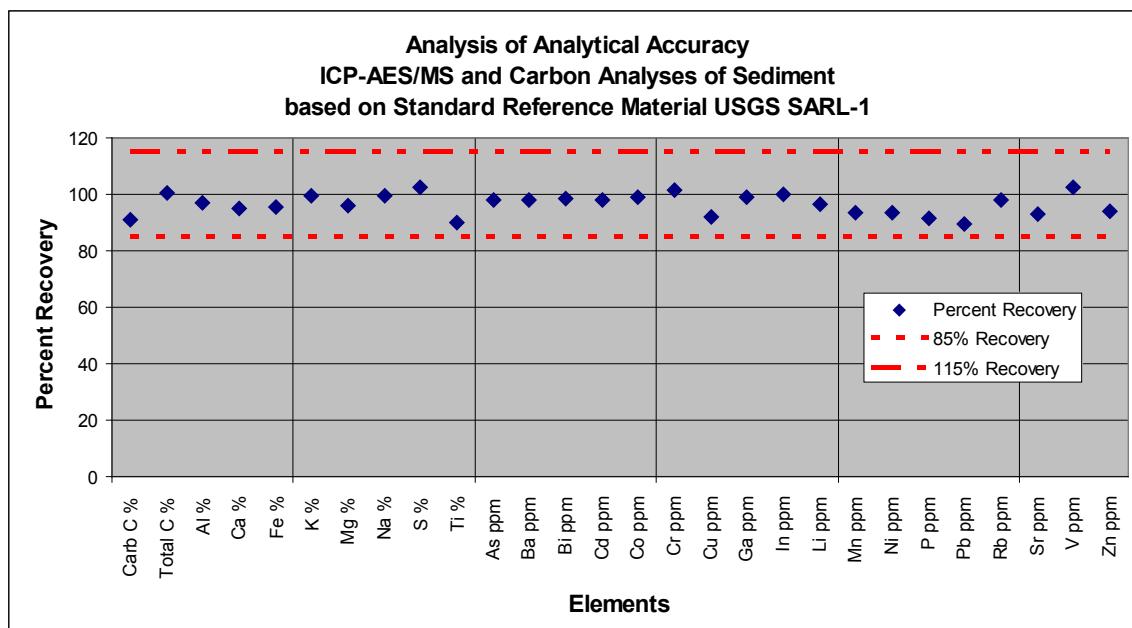


Figure 2–8. Accuracy plot for ICP-AES/MS and carbon analyses for reference material USGS SAR-L. [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; USGS, U.S. Geological Survey; SAR-L, sediment, Animas River watershed, Colo., low levels of contaminants; %, percent; ppm, parts per million]

Table 2–8. Summary statistics for assessing analytical variation and accuracy for the reference material USGS SAR-L determined after a four-acid total digestion of sediment samples by the ICP-AES/MS package at SGS Minerals and for total and carbonate carbon. ([Click to see Excel file.](#))

[USGS, U.S. Geological Survey; SAR-L, sediment, Animas River watershed, Colo., low levels of contaminants; ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; RL, reporting limit; n, number of analyses of SAR-L; RSD, relative standard deviation; %, percent; ppm, parts per million]

Element	Units	RL	n	Target value	Mean	Standard deviation	% RSD	% Recovery
Al	%	0.01	7	5.84	5.66	0.15	2.7	97
Ca	%	0.01	7	1.06	1.01	0.016	1.56	95
Fe	%	0.01	7	2.60	2.48	0.071	2.85	95
K	%	0.01	7	3.02	3.01	0.084	2.78	100
Mg	%	0.01	7	0.54	0.514	0.014	2.72	96
Na	%	0.01	7	1.51	1.50	0.041	2.76	99
S	%	0.01	7	0.08	0.080	0.000	0.00	103
Ti	%	0.01	7	0.23	0.207	0.0049	2.36	90
As	ppm	1	7	16.3	16.0	0.0	0.0	98
Ba	ppm	5	7	910	893	18.4	2.06	98
Bi	ppm	0.04	7	1.19	1.17	0.23	19.4	99
Cd	ppm	0.1	7	2.95	2.89	0.15	5.07	98
Co	ppm	0.1	7	7.14	7.09	0.27	3.77	99
Cr	ppm	1	7	96	97	3.9	4.01	101
Cu	ppm	0.5	7	386	356	14.8	4.16	92
Ga	ppm	0.05	7	16.5	16.4	0.72	4.39	99
In	ppm	0.02	7	0.30	0.30	0.01	3.83	100
Li	ppm	1	7	26.4	25	0.98	3.84	96
Mn	ppm	5	7	2,190	2,046	70.4	3.44	93
Ni	ppm	0.5	7	53	49.5	1.38	2.80	93
P	ppm	50	7	830	759	9.00	1.19	91
Pb	ppm	0.5	7	589	527	22.5	4.27	89
Rb	ppm	0.2	7	140	137	6.61	4.82	98
Sr	ppm	0.5	7	156	145	2.89	1.99	93
V	ppm	1	7	137	140	3.82	2.72	102
Zn	ppm	1	7	457	431	5.91	1.37	94
Total C	%	0.01	7	1.06	1.07	0.01	0.89	101
Carbonate C	%	0.003	7	0.11	0.01	0.0	0.0	91

Reference Material USGS SAR-M

Like USGS GSP-QC and SAR-L, the reference material SAR-M is not certified nor used outside of the U.S. Geological Survey. It is a composite of contaminated sediment from the Animas River watershed in Colorado with moderate levels of contaminant constituents. This material was developed in-house at the Denver USGS laboratories for inclusion with samples sent to the SGS Minerals laboratory. During the period of September 2004 to November 2008, this material was analyzed 320 times by the contract laboratory for the ICP-AES/MS package as blind samples and 250 times for carbon. The results were compiled, and the means derived were used as target values for the accuracy assessments of this material submitted in this report. As a consequence, target values are available for all elements in the ICP-AES/MS package and for total and carbonate carbon.

The noncertified USGS reference material SAR-M was analyzed six times along with the study samples. Table 2–9 gives the percent RSD and percent recovery results for this standard, and figure 2–9 shows graphically the accuracy plot. All of the elements plot within ± 15 percent for percent recovery, except for vanadium, which had a 170 percent recovery.

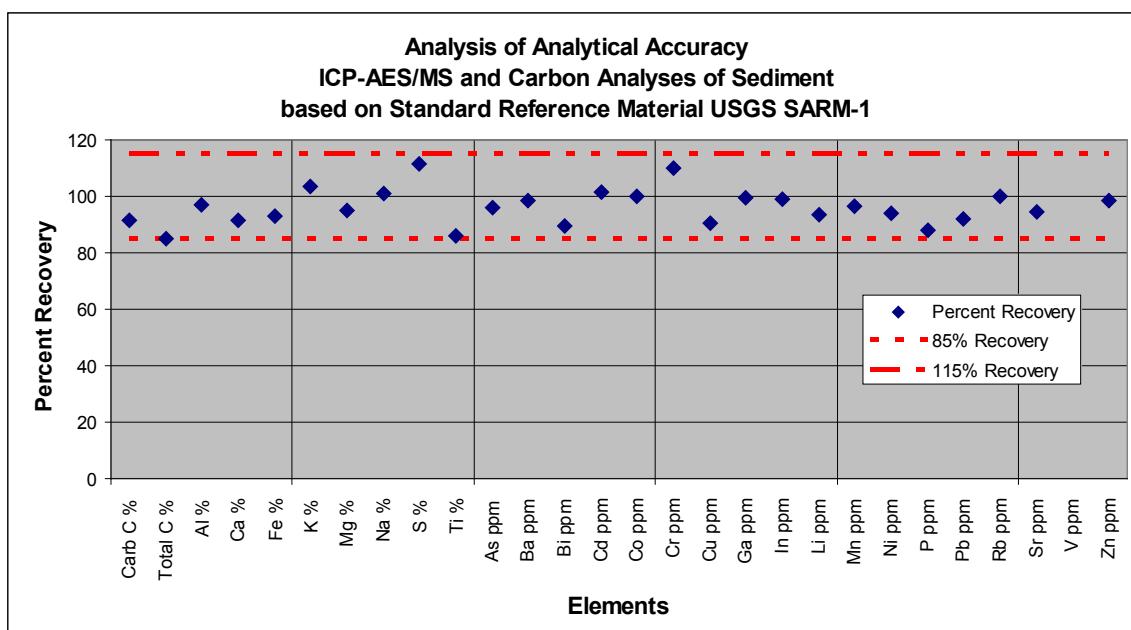


Figure 2–9. Accuracy plot for ICP-AES/MS and carbon analyses for reference material USGS SAR-M. [ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; USGS, U.S. Geological Survey; SAR-M, sediment, Animas River watershed, Colo., moderate levels of contaminants; %, percent; ppm, parts per million]

Table 2–9. Summary statistics for assessing analytical variation and accuracy for the reference material USGS SAR-M determined after a four-acid total digestion of sediment samples by the ICP-AES/MS package at SGS Minerals and for total and carbonate carbon. ([Click to see Excel file.](#))

[USGS, U.S. Geological Survey; SAR-M, sediment, Animas River watershed, Colo., moderate levels of contaminants; ICP-AES/MS, inductively coupled plasma–atomic emission spectroscopy/mass spectroscopy; RL, reporting limit; n, number of analyses of SAR-M; RSD, relative standard deviation; %, percent; ppm, parts per million]

Element	Units	RL	n	Target value	Mean	Standard deviation	% RSD	% Recovery
Al	%	0.01	6	6.13	5.94	0.10	1.7	97
Ca	%	0.01	6	0.587	0.55	0.02	3.75	93
Fe	%	0.01	6	3.1	2.92	0.09	3.14	94
K	%	0.01	6	2.88	2.97	0.09	3.20	103
Mg	%	0.01	6	0.49	0.47	0.01	3.01	96
Na	%	0.01	6	1.16	1.17	0.02	1.67	101
S	%	0.01	6	0.118	0.13	0.01	6.63	110
Ti	%	0.01	6	0.272	0.24	0.02	8.40	88
As	ppm	1	6	38.8	37.5	1.09	2.91	97
Ba	ppm	5	6	801	790	8.82	1.12	99
Bi	ppm	0.04	6	1.94	1.77	0.16	9.05	91
Cd	ppm	0.1	6	5.27	5.34	0.10	1.88	101
Co	ppm	0.1	6	10.7	10.7	0.28	2.59	100
Cr	ppm	1	6	79.7	86.6	4.15	4.79	109
Cu	ppm	0.5	6	331	304.	15.9	5.23	92
Ga	ppm	0.05	6	16.8	16.7	0.95	5.70	99
In	ppm	0.02	6	1.08	1.07	0.04	3.35	99
Li	ppm	1	6	27.4	25.9	1.29	4.96	95
Mn	ppm	5	6	5,220	5,068	152	3.00	97
Ni	ppm	0.5	6	41.5	39.3	1.37	3.48	95
P	ppm	50	6	775	696	37.3	5.35	90
Pb	ppm	0.5	6	982	915	42.4	4.64	93
Rb	ppm	0.2	6	146	146	6.04	4.14	100
Sr	ppm	0.5	6	151	144	3.32	2.30	95
V	ppm	1	6	37.2	63.5	11.6	18.3	171
Zn	ppm	1	6	930	919	24.3	2.64	99
Total C	%	0.01	6	0.35	0.30	0.03	9.46	87
Carbonate C	%	0.003	6	0.13	0.13	0.02	15.4	100